

PATENT APPLICATION

INHIBITORS OF CATHEPSIN S

Inventors:

Hong Liu, a citizen of P. R. China., residing at 6348 Oleander Way, San Diego, CA 92130

David Tully, a citizen of the United States of America, residing at San Diego, California

Robert Epple, a citizen of Austria, residing at San Diego, California

Badry Bursulaya, a citizen of Russian Federation, residing at San Diego, California

Jennifer Williams, a citizen of the United Kingdom, residing at San Diego, California

Arnab Chatterjee, a citizen of the United States of America, residing at Encinitas, California

Jennifer Leslie Harris, a citizen of the United States of America, residing at San Diego, California

Jun Li, a citizen of the People's Republic of China, residing at San Diego, California

Assignee: IRM LLC
Sophia House, 48 Church Street
P. O. Box HM 2899
HM LX Hamilton
Bermuda

Entity: Large

TOWNSEND and TOWNSEND and CREW LLP
Two Embarcadero Center, 8th Floor
San Francisco, California 94111-3834
Tel: 925-472-5000

INHIBITORS OF CATHEPSIN S

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] The present application claims priority to U.S. Patent No. 60/457,595, filed March 24, 2003, the teachings of which are hereby incorporated by reference in their entirety.

5

BACKGROUND OF THE INVENTION

[0002] Cysteine proteases represent an enzymatic class of proteins that catalyze the hydrolysis of peptide bonds by a nucleophilic sulfhydryl group of a cysteine residue in the active site of the enzyme. Several normal and disease processes in mammals have been associated with cysteine protease activity and include, but are not limited to: osteoporosis, osteoarthritis (Inui, T., O. Ishibashi, *J Biol Chem* **1997**, *272*(13), 8109-12; Saftig, P., E. Hunziker, *et al.*, *Adv Exp Med Biol* **2000**+*ADs* **2000**, *477*, 293-303; Saftig, P., E. Hunziker, *et al.*, *Proc Natl Acad Sci U S A* **1998**, *95*(23), 13453-8), periodontal diseases, Paget's disease, atherosclerosis (Jormsjo, S., D. M. Wuttge, *et al.*, *Am J Pathol* **2002** *161*(3), 939-45), multiple sclerosis (Beck, H., G. Schwarz, *et al.*, *Eur J Immunol* **2001**, *31*(12), 3726-36),
10 rheumatoid arthritis (Nakagawa, T. Y., W. H. Brissette, *et al.*, *Immunity* **1999**, *10*(2), 207-17; Hou, W. S., Z. Li, *et al.*, *Am J Pathol* **2001**, *159*(6), 2167-77), juvenile onset diabetes, lupus, asthma (Cimerman, N., P. M. Brguljan, *et al.*, *Pflugers Arch* **2001**, *442*(6 Suppl 1), R204-6), tissue rejection, Alzheimer's disease (Lemere, C. A., J. S. Munger, *et al.*, *Am J Pathol* **1995**, *146*(4), 848-60), Parkinson's disease (Liu, Y., L. Fallon, *et al.*, *Cell* **2002**, *111*(2), 209-18),
15 neuronal degeneration, shock (Jaeschke, H., M. A. Fisher, *et al.*, *J Immunol* **1998**, *160*(7), 3480-6), cancer (Fernandez, P. L., X. Farre, *et al.*, *Int J Cancer* **2001**, *95*(1), 51-5), malaria (Malhotra, P., P. V. Dasaradhi, *et al.*, *Mol Microbiol* **2002**, *45*(5), 1245-54), Chagas (Eakin, A. E., A. A. Mills, *et al.*, *J Biol Chem* **1992**, *267*(11), 7411-20), leishmaniasis, shistosomiasis, and African trypanosomiasis (Caffrey, C. R., S. Scory, *et al.*, *Curr Drug Targets* **2000**, *1*(2),
20 155-62; Lalmanach, G., A. Boulange, *et al.*, *Biol Chem* **2002**, *383*(5), 739-49).
25

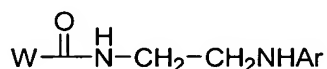
[0003] Cathepsins are a subclass of cysteine protease that belong to the enzyme classification EC 3.4.22 (Barrett, A. J., N. D. Rawlings, *et al.*, *Handbook of proteolytic enzymes*. London, Academic Press). Cathepsins play a major role in lysosomal, endosomal, and extracellular protein degradation and have thus been implicated in many disease
30 processes. For example, Cathepsin B [EC 3.4.22.1] has been postulated to play a role in tumor metastasis (Berquin, I. M. and B. F. Sloane *Adv Exp Med Biol* **1996**, *389*, 281-94).

[0004] Cathepsin S [EC 3.4.22.27] is largely expressed in professional antigen presenting cells such as macrophages and dendritic cells. Cathepsin S has been shown to be required for proper MHC class II antigen presentation (Shi, G. P., J. A. Villadangos, *et al.*, *Immunity* 1999, 10(2) 197-206). As a result of its non-redundant role in MHC class II antigen presentation, cathepsin S has been associated with inflammation, arthritis, and atherosclerosis. The selective expression of cathepsin K [EC 3.4.22.38] in osteoclasts coupled with the ability of cathepsin K to degrade type I collagen suggests that it plays a role in normal and pathogenic bone remodeling (Bromme, D., K. Okamoto, *et al.*, *J Biol Chem* 1996, 271(4), 2126-32). There is a need in the art for compounds and methods that selectively inhibit specific cysteine proteases for treating several pathogenic disorders in mammals. The present invention satisfies these and other needs.

SUMMARY OF THE INVENTION

[0005] The present invention provides compounds, compositions and methods for the selective inhibition of cathepsin S. The compounds of the present invention are selective for cathepsin S in the presence of other cathepsin isozymes (*e.g.*, cathepsin K). In a preferred embodiment, the compounds of the present invention are selective for cathepsin S in the presence of cathepsin K. The present invention also provides methods for treating a disease state in a subject by selectively inhibiting cathepsin S in the presence of other cathepsin isozymes. In a preferred aspect, cathepsin S is selectively inhibited in the presence of cathepsin K.

[0006] As such, the present invention provides a compound of Formula I:



(I)

or a pharmaceutically acceptable salt or prodrug thereof,

wherein:

W is a member selected from the group of :

$\text{R}^1\text{-X-(C=O)-NH-CHR}^2\text{-}$,

$\text{R}^4\text{-Y-(C=O)-NH-CHR}^3\text{-}$,

$\text{R}^6\text{-(C=O)-NH-CHR}^5\text{-}$,

$\text{R}^7\text{-NH-(C=O)-NH-CHR}^8\text{-}$,

$\text{R}^{10}\text{-Z-(C=O)-NH-CHR}^9\text{-}$, and

$R^{11}-(C=O)-NH-CHR^{12}-$;

R^1 is a member selected from the group of phenyl substituted with 0-2 R^{1a} , pyridyl substituted with 0-2 R^{1a} , and pyridinium N-oxide substituted with 0-2 R^{1a} ; each R^{1a} is independently a member selected from the group of Cl, F, OCF_3 , OCH_3 , CH_3 and CF_3 ;

X is a member selected from the group of furanylene substituted with 0-1 R^x , thienylene substituted with 0-1 R^x , pyrazolylene substituted with 0-1 R^x , thiazolylene substituted with 0-1 R^x , and oxazolylene substituted with 0-1 R^x ;

R^x is a member selected from the group of F, Cl, CH_3 and CF_3 ;

R^2 is a member selected from the group of phenyl substituted with 0-2 R^{2a} , and $(CH_2)_nR^{2b}$;

each R^{2a} is independently a member selected from the group of Cl, F, OCF_3 , OCH_3 , CH_3 and CF_3 ;

R^{2b} is independently a member selected from the group of phenyl substituted with 0-2 R^{2a} ; cyclopentyl, cyclohexyl and tetrahydropyranyl;

n is the integer 1 or 2;

R^3 is $(CH_2)_mR^{3b}$;

R^{3b} is selected from the group of phenyl substituted with 0-2 R^{2a} , cyclopentyl and cyclohexyl;

m is the integer 1 or 2;

R^4 is a member selected from the group of phenyl substituted with 0-3 R^{4a} , thienyl, tetrazolyl, cyclopentenyl and indolyl;

each R^{4a} is a member selected from the group of phenyl, OH, C_1 - C_4 alkyl, C_1 - C_4 alkoxy, CF_3 , OCF_3 , F, Cl, $CH_3S(=O)_2-$, morpholinyl, pyrrolidinyl, piperidinyl and 4-acetylpiperazinyl;

Y is a member selected from the group of $-CR^{17}R^{18}$, $-NH-CH_2-$ and $-O-CH_2-$;

R^5 is a member selected from the group of phenyl substituted with 0-2 R^{5a} , thiophene, naphthyl, and CH_2R^{5b} , CH_2CH_2 (cyclohexyl), $CH_2CH_2CH_2$ (cyclohexyl), CH_2CH_2Ph , $CH(CH_3)R^{5c}$, $CH_2CH=CHPh$, $-CH_2OCH_2Ph$, $-CH(CH_3)OCH_2Ph$; each R^{5a} is independently a member selected from the group of F, Cl, NO_2 , OCH_3 , OCH_2Ph , OPh , CH_3 , OCF_3 and CF_3 ;

R^{5b} is independently a member selected from the group of phenyl substituted with 0-2 R^{5c} ; cyclopentyl, cyclohexyl, naphthyl, indolyl and pyridyl;

R^{5c} is independently a member selected from the group of OH, Cl, F, Br, I, CN, NO₂, CH₃, OCH₃, ^tBu, O-^tBu, -NHC(=O)CH₃, CF₃, OCF₃; phenyl substituted with 0-2 R^{5d} ; phenoxy substituted with 0-2 R^{5d} ; benzyloxy substituted with 0-2 R^{5d} ; pyridyl substituted with 0-2 R^{5d} ; pyrimidinyl substituted with 0-2 R^{5d} ; thienyl substituted with 0-2 R^{5d} ;

R^{5d} is independently a member selected from the group of CH₃, Cl, F, OCH₃, CF₃, OCF₃, N(CH₃)₂, acetyl, OH, CH₂OH, NH₂, CN and NO₂;

R^{5e} is phenyl substituted with 0-2 R^{5a} ;

R^6 is a member selected from the group of phenyl substituted with 0-3 R^{6a} , furanyl substituted with 0-2 R^{6b} , thienyl substituted with 0-2 R^{6b} , oxazolyl substituted with 0-2 R^{6b} , thiazolyl substituted with 0-2 R^{6b} , pyridyl, pyridazinyl and cyclopropyl;

each R^{6a} is independently a member selected from the group of Cl, F, Br, OCF₃, CF₃, C₁-C₄ alkyl, C₁-C₄ alkoxy, -S(=O)₂CH₃, CN, -N(CH₃)₂, OCF₂H, -CH₂-benzimidazole, -NH-S(=O)₂CH₃, -NR¹³R¹⁴, OR¹⁴, CH₂-morpholine, CH₂NH₂, OCH₂Ph, and OPh;

alternatively, two R^{6a} substituents on adjacent atoms may be combined to form a 5 to 6 membered heterocyclic fused radical, wherein said 5 to 6 membered heterocyclic fused radical has 1 or 2 oxygen atom(s);

each R^{6b} is independently a member selected from the group of NH₂, F, Cl, Br, -S(=O)₂R¹⁵, CH₃, and CF₃;

R^7 is a member selected from the group of (CH₂)_p R^{7a}, and naphthyl substituted with 0-2 R^{7b} ;

p is the integer 0, 1, or 2;

R^{7a} is phenyl substituted with 0-2 R^{7b} ;

R^{7b} is a member selected from the group of F, Cl, CF₃, C₁-C₄ alkyl, C₁-C₄ alkoxy, OCF₃, phenoxy and acetyl;

alternatively, two R^{7b} substituents on adjacent atoms may be combined to form a 5 to 6 membered heterocyclic fused radical, wherein said 5 to 6 membered heterocyclic fused radical has 1 or 2 oxygen atom(s);

R^8 is -CH₂-R^{3b};

R^9 is (CH₂)_qR^{9a};

R^{9a} is a member selected from the group of cyclopentyl, phenyl and cyclohexyl;

q is the integer 1 or 2;

R¹⁰ is a member selected from the group of phenyl substituted with 0-2 R^{10a}, 5
membered heteroaryl containing 1 to 4 heteroatoms each independently a
member selected from the group of N, O and S, wherein said heteroaryl is
substituted with 0-2 R^{10a}, 6 membered heteroaryl containing 1 to 2 N, wherein
5 said heteroaryl is substituted with 0-2 R^{10a}, morpholinyl substituted with 0-2
R^{10a}, piperazinyl substituted with 0-2 R^{10a} and piperidinyl substituted with 0-2
R^{10a};

each R^{10a} is independently a member selected from the group of Cl, F, C₁-C₄ alkyl,
C₁-C₄ alkoxy, OCF₃, and CF₃;

10 alternatively, two R^{10a} substituents on adjacent atoms may be combined to form a 5 to
6 membered heterocyclic fused radical, wherein said 5 to 6 membered
heterocyclic fused radical comprises 1 or 2 heteroatom(s);

Z is phenylene;

R¹¹ is a member selected from the group of indolyl substituted with 0-2 R^{11a},
15 benzofuranyl substituted with 0-2 R^{11a}, benzothienyl substituted with 0-2 R^{11a},
benzothiazole substituted with 0-2 R^{11a}, benzisoxazolyl substituted with 0-2
R^{11a}, benzoxazolyl substituted with 0-2 R^{11a}, and pyrazolo[1,5-a]pyrimidinyl
substituted with 0-2 R^{11a}, piperidinyl N-substituted with 0-1 R^{11b}, morpholinyl
N-substituted with 0-1 R^{11b}; and 2-oxo-pyrrolidinyl with 0-1 R^{11b};

20 each R^{11a} is independently a member selected from the group of Cl, F, NH₂, CH₃,
OCH₃, -C(=O)OCH₃, OCF₃, and CF₃;

each R^{11b} is independently a member selected from the group of pyrimidinyl
substituted with 0-2 R^{11c}; benzyl, acetyl, CH₂-furanyl, and CH₂-thienyl;

each R^{11c} is independently a member selected from the group of Br and CH₃;

25 R¹² is (CH₂)_sR^{12a};

R^{12a} is a member selected from the group of cyclopentyl and cyclohexyl;

s is the integer 1 or 2;

R¹³ is a member selected from the group of H and C₁-C₄ alkyl;

R¹⁴ is pyrimidinyl substituted with 0-2 R¹⁶;

30 R¹⁵ is a member selected from the group of C₁-C₄ alkyl, morpholinyl, pyrrolidinyl and
piperidinyl;

R¹⁶ is a member selected from the group of CH₃ and OCH₃;

each of R¹⁷ and R¹⁸ is independently a member of H, OH, F, phenyl and C₁-C₃ alkyl;
alternatively, R¹⁷ and R¹⁸ may be taken together to form a C₃-C₆ cycloalkyl;

Ar is a phenyl substituted with 0-2 R¹⁹; and
each R¹⁹ is independently a member selected from the group consisting of F, Cl, COOH,
C₁-C₄ alkoxy, OCHF₂ and OCF₃.

5 [0007] In a second aspect, the present invention provides a pharmaceutical composition
comprising a compound of Formula I, as described above, and a pharmaceutically acceptable
excipient.

[0008] In a third aspect, the present invention provides a method of selectively inhibiting
the cathepsin S activity in a mammal in need thereof, comprising administering to the
mammal a therapeutically effective amount of a compound of Formula I, as described above,
10 or a pharmaceutically acceptable salt or prodrug thereof.

[0009] These and other aspects, objects and embodiments will become more apparent when
read with the accompanying figure and detailed description which follows.

DESCRIPTION OF THE DRAWINGS

15 [0010] Figure 1 depicts MHC II antigen presentation.

DETAILED DESCRIPTION OF THE INVENTION

I. Definitions

20 [0011] Unless defined otherwise, all technical and scientific terms used herein generally
have the same meaning as commonly understood by one of ordinary skill in the art to which
this invention belongs. Generally, the nomenclature used herein and the laboratory
procedures for organic and analytical chemistry are those well known and commonly
employed in the art.

[0012] As used in this disclosure, the following abbreviations and terms have the defined
25 meaning, unless expressly modified in the context in which the term is used:

Ac	acetyl
Bn	benzyl
Boc	t-butoxycarbonyl
Cbz or Z	benzyloxycarbonyl

	DCC	N,N'-dicyclohexylcarbodiimide
	DCM	dichloromethane
	DIBAL	diisobutylaluminum hydride
	DIC	N,N'-diisopropylcarbodiimide
5	DIEA or DIPEA	diisopropylethylamine
	DMAP	4-(dimethylamino)pyridine
	DMF	dimethylformamide
	DMSO	dimethyl sulfoxide
	EDC or EDCI	1-ethyl-3-(dimethylaminopropyl)-carbodiimide
10	Fmoc	9-fluorenylmethoxycarbonyl
	HATU	O-(7-azabenzotriazol-1-yl)-N,N,N',N'-tetramethyluronium hexafluorophosphate
	HOBt	1-hydroxybenzotriazole
	KHMDS	potassium hexamethyldisilazide
15	LAH	lithium aluminum hydride
	LDA	lithium diisopropylamide
	LHMDS	lithium hexamethyldisilazide
	m-CPBA	m-chloroperbenzoic acid
	MW	microwave
20	NaHMDS	sodium hexamethyldisilazide
	PCC	pyridinium chlorochromate
	PDC	pyridinium dichromate
	PG	protecting group
	PTSA	p-toluenesulfonic acid
25	Py	pyridine
	RT or rt	room temperature
	TEA	triethylamine
	Tf	trifluoromethanesulfonyl
	TFA	trifluoroacetic acid
30	THF	tetrahydrofuran
	Tol	p-tolyl
	TPAP	tetrapropylammonium perruthenate

[0013] The term “lower” referred to above and hereinafter in connection with organic radicals or compounds respectively defines a compound or radical which can be branched or

unbranched with up to and including 7, preferably up to and including 4 and (as unbranched) one or two carbon atoms.

5 [0014] The term “perfluoro” referred to above and hereinafter in connection with organic radicals or compounds respectively, defines a compound or radical which has at least two available hydrogens substituted with fluorine. For example, perfluorophenyl refers to 1,2,3,4,5-pentafluorophenyl, perfluoromethyl refers to 1,1,1-trifluoromethyl, and perfluoromethoxy refers to 1,1,1-trifluoromethoxy.

10 [0015] An alkyl group is branched or unbranched and contains 1 to 7 carbon atoms, preferably 1-4 carbon atoms. Alkyl represents, for example, methyl, ethyl, propyl, butyl, isopropyl or isobutyl.

[0016] Alkenyl represents either straight chain or branched alkenyl of 2 to 7 carbon atoms, preferably 2-4 carbon atoms, e.g. as vinyl, propenyl, isopropenyl, butenyl, isobutenyl or butadienyl.

15 [0017] Alkynyl represents either straight chain or branched alkynyl of 2 to 7 carbon atoms, preferably 2-4 carbon atoms, e.g. as acetylenyl, propynyl, isoprpropynyl, butynyl or isobutynyl.

[0018] Alkyl, alkenyl or alkynyl can be substituted by up to 3 substituents selected from alkoxy, aryl, heterocyclyl, hydroxy, halogen, cyano, optionally substituted amino, or optionally substituted amino-oxy or trifluoromethyl.

20 [0019] Alkylene represents either straight chain or branched alkylene of 1 to 7 carbon atoms, i.e. a divalent hydrocarbon radical of 1 to 7 carbon atoms; for instance, straight chain alkylene being the bivalent radical of Formula $-(CH_2)_n$, where n is 1, 2, 3, 4, 5, 6 or 7. Preferably alkylene represents straight chain alkylene of 1 to 4 carbon atoms, e.g. a methylene, ethylene, propylene or butylene chain, or the methylene, ethylene, propylene or butylene chain mono-substituted by C₁-C₃-alkyl (preferably methyl) or disubstituted on the
25 same or different carbon atoms by C₁-C₃-alkyl (preferably methyl), the total number of carbon atoms being up to and including 7.

[0020] An alkoxy (or alkyloxy) group preferably contains 1-7 carbon atoms, more preferably 1-6 carbon atoms, and represents for example ethoxy, propoxy, isopropoxy, isobutoxy, preferably methoxy. Alkoxy includes cycloalkyloxy and cycloalkyl-alkyloxy.
30

[0021] Halogen (halo) preferably represents chloro or fluoro, but may also be bromo or iodo.

[0022] Aryl represents monocyclic, bicyclic or tricyclic aryl, for example, phenyl or phenyl mono-, di- or tri-substituted by one, two or three radicals selected from alkyl, alkoxy, aryl, hydroxy, halogen, cyano, amino, amino-alkyl, trifluoromethyl, alkylenedioxy and oxy-C₂-C₃-alkylene; all of which are optionally further substituted, for instance as hereinbefore defined; or 1- or 2-naphthyl; or 1- or 2-phenanthrenyl. Alkylenedioxy is a divalent substitute attached to two adjacent carbon atoms of phenyl, e.g. methylenedioxy or ethylenedioxy. Oxy-C₂-C₃-alkylene is also a divalent substituent attached to two adjacent carbon atoms of phenyl, e.g. oxyethylene or oxypropylene. An example for oxy- C₂-C₃-alkylene-phenyl is 2,3-dihydrobenzofuran-5-yl.

[0023] Preferred as aryl is naphthyl, phenyl or phenyl mono- or disubstituted by alkoxy, phenyl, halogen, alkyl or trifluoromethyl, especially phenyl or phenyl-mono- or disubstituted by alkoxy, halogen or trifluoromethyl, and in particular phenyl.

[0024] Examples of substituted phenyl groups as R are, e.g. 4-chlorophen-1-yl, 3,4-dichlorophen-1-yl, 4-methoxyphen-1-yl, 4-methylphen-1-yl, 4-aminomethylphen-1-yl, 4-methoxyethylaminomethylphen-1-yl, 4-hydroxyethylaminomethylphen-1-yl, 4-hydroxyethyl-(methyl)-aminomethylphen-1-yl, 3-aminomethylphen-1-yl, 4-N-acetylaminoethylphen-1-yl, 4-aminophen-1-yl, 3-aminophen-1-yl, 2-aminophen-1-yl, 4-phenyl-phen-1-yl, 4-(imidazol-1-yl)-phen-yl, 4-(imidazol-1-ylmethyl)-phen-1-yl, 4-(morpholin-1-yl)-phen-1-yl, 4-(morpholin-1-ylmethyl)-phen-1-yl, 4-(2-methoxyethylaminomethyl)-phen-1-yl and 4-(pyrrolidin-1-ylmethyl)-phen-1-yl, 4-(thiophenyl)-phen-1-yl, 4-(3-thiophenyl)-phen-1-yl, 4-(4-methylpiperazin-1-yl)-phen-1-yl, and 4-(piperidiny)-phenyl and 4-(pyridiny)-phenyl optionally substituted in the heterocyclic ring.

[0025] Benzyl represents a phenyl-CH₂ - group. Substituted benzyl means a benzyl group in which the phenyl ring is substituted with one or more ring system substituents. Representative benzyl groups include 4-bromobenzyl, 4-methoxybenzyl, 2,4-dimethoxybenzyl, and the like.

[0026] Heteroaryl represents monocyclic or bicyclic heteroaryl, for example pyridyl, pyridyl N-oxide, indolyl, indazolyl, quinoxaliny, quinoliny, isoquinoliny, benzothienyl, benzofuranyl, benzopyranyl, benzothiopyranyl, furanyl, pyrrolyl, thiazolyl, benzothiazolyl, oxazolyl, isoxazolyl, triazolyl, tetrazolyl, pyrazolyl, imidazolyl, thienyl, or any other radicals

substituted, especially mono- or di-substituted, by e.g. alkyl, nitro or halogen. Pyridyl represents 2-, 3- or 4-pyridyl, advantageously 2- or 3-pyridyl. Thienyl represents 2- or 3-thienyl. Quinoliny represents preferably 2-, 3- or 4-quinoliny. Isoquinoliny represents preferably 1-, 3- or 4-isoquinoliny. Benzopyranyl, benzothiopyranyl represents preferably 3-benzopyranyl or 3-benzothiopyranyl, respectively. Thiazolyl represents preferably 2- or 4-thiazolyl, and most preferred, 4-thiazolyl. Triazolyl is preferably 1-, 2- or 5-(1,2,4-triazolyl). Tetrazolyl is preferably 5-tetrazolyl.

[0027] Preferably, heteroaryl is pyridyl, pyridyl N-oxide, indolyl, quinoliny, pyrrolyl, thiazolyl, isoxazolyl, triazolyl, tetrazolyl, pyrazolyl, imidazolyl, thienyl, furanyl, benzothiazolyl, benzofuranyl, isoquinoliny, benzothienyl, oxazolyl, indazolyl, or any of the radicals substituted, especially mono- or di-substituted.

[0028] Biaryl may preferably be, e.g., biphenyl, namely 2, 3 or 4-biphenyl, preferably, 4-biphenyl, each optionally substituted by, e.g., alkyl, alkoxy, halogen, trifluoromethyl or cyano, or heterocyclic-carbocyclic biaryl, preferably, e.g., thienylphenyl, pyrrolylphenyl and pyrazolylphenyl.

[0029] Cycloalkyl represents a saturated cyclic hydrocarbon optionally substituted by alkyl which contains 3 to 10 ring carbons and is advantageously cyclopentyl, cyclohexyl, cycloheptyl or cyclooctyl optionally substituted by alkyl.

[0030] Bicycloalkyl represents a saturated bicyclic ring group of 7-15 carbon atoms.

Exemplary bicycloalkyl rings include [3.3.0]bicyclooctanyl, [2.2.2]bicyclooctanyl, [4.3.0]bicyclononane, [4.4.0]bicyclodecane (decalin), spiro[3.4]octanyl, spiro[2.5]octanyl, and so forth, optionally substituted by alkyl.

[0031] Amino can be optionally substituted by, e.g., alkyl.

[0032] Carbocyclic represents a saturated or partially unsaturated cyclic hydrocarbon with 5 to 7 ring members, wherein 1 to 2 ring members can optionally be replaced with one of the following groups: -O-, -S-, -S(=O)-, -S(=O)₂- and -NR-, wherein R is a radical of the present invention.

[0033] Heterocyclyl represents a saturated cyclic hydrocarbon containing one or more, preferably 1 or 2 heteroatoms selected from O, N or S, and from 3 to 10, preferably 5 to 8, ring atoms; for example, tetrahydrofuranyl, tetrahydrothienyl, tetrahydropyrrolyl, piperidinyl,

piperazinyl or morpholino; all of which can be optionally substituted, for instance as hereinbefore defined for aryl.

[0034] Pharmaceutically acceptable salts of the acidic compounds of the present invention are salts formed with bases, namely cationic salts such as alkali and alkaline earth metal salts, such as sodium, lithium, potassium, calcium, magnesium, as well as ammonium salts, such as ammonium, trimethyl-ammonium, diethylammonium, and tris-(hydroxymethyl)-methyl-ammonium salts.

[0035] Similarly acid addition salts, such as of mineral acids, organic carboxylic and organic sulfonic acids, e.g., hydrochloric acid, methanesulfonic acid, maleic acid, are also possible provided a basic group, such as pyridyl, constitutes part of the structure.

[0036] "Treat", "treating" and "treatment" refer to a method of alleviating or abating a disease and/or its attendant symptoms.

[0037] "Inhibition", "inhibits" and "inhibitor" refer to a compound that prohibits, or a method of prohibiting, a specific action or function.

[0038] "Inhibition constant", K_i , is the dissociation constant of the enzyme-inhibitor complex, or the reciprocal of the binding affinity of the inhibitor to the enzyme. For classical inhibition the value of K_i is much greater than the enzyme concentration and the K_i can be measured by monitoring the rate of reaction for a competitive substrate at multiple inhibitor concentrations. The inhibited rates are then fit by nonlinear regression to the following equation:

$$v_i/v_o = \frac{K_m + [S]}{K_m (1 + [I]/K_i) + [S]}$$

where v_o is the initial rate of substrate processing in the absence of inhibitor, v_i is the initial rate of substrate processing at a concentration $[I]$ of inhibitor, K_m is the steady state Michaelis constant (Fersht, A. Structure and Mechanism in Protein Science. New York, W.H. Freeman and Company, 1999), and $[S]$ is the concentration of competitive substrate.

[0039] The assumption being made for the classical inhibition described above is that the free inhibitor concentration is equal to the total inhibitor concentration. For inhibitors that have K_i 's that are approximately equal to the enzyme concentration $[E]$, the assumption that the free inhibitor concentration is equal to the total inhibitor concentration is no longer valid

and an alternative equation has to be fit for determination of the apparent inhibition constant, K_i^{app} using described methods (Kuzmic, P., K. C. Elrod, *et al.*, *Anal Biochem* **2000**, 286(1), 45-50):

$$v_i/v_o = \frac{[E]-[I]-K_i^{app} + \text{SQRT} \left(([E]-[I]-K_i^{app})^2 + 4[E] K_i^{app} \right)}{2[E]}$$

- 5 The inhibition constant, K_i , can be determined from the apparent inhibition constant, K_i^{app} , for competitive inhibitors by using the following relationship:

$$K_i = \frac{K_i^{app}}{1 + [S]/K_m}$$

10 [0040] “Therapeutically effective amount” refers to that amount of the compound being administered sufficient to prevent development of or alleviate to some extent one or more of the symptoms of the condition or disorder being treated.

15 [0041] “Composition” as used herein is intended to encompass a product comprising the specified ingredients in the specified amounts, as well as any product which results, directly or indirectly, from combination of the specified ingredients in the specified amounts. By “pharmaceutically acceptable” it is meant the carrier, diluent or excipient must be compatible with the other ingredients of the Formulation and deleterious to the recipient thereof.

[0042] “Subject” refers to animals such as mammals, including, but not limited to, primates (*e.g.*, humans), cows, sheep, goats, horses, dogs, cats, rabbits, rats, mice and the like. In certain aspects, the subject is a human.

20 [0043] “Prodrug” refers to the compounds of this invention which may be modified by appending appropriate functionalities to enhance selective biological properties. Such modifications are known in the art and include those which increase penetration into a given biological compartment (*e.g.* central nervous system), increase oral bioavailability, increase solubility to allow administration by injection, alter metabolism and alter rate and/or route of excretion. In addition, the compounds may be altered to prodrug form such that the desired
25 compound is created in the body of the patient as the result of the action of metabolic or other biochemical processes on the prodrug.

[0044] It will be apparent to one skilled in the art that certain compounds of this invention may exist in tautomeric forms or hydrated forms, all such forms of the compounds being within the scope of the invention.

[0045] Structures depicted herein are also meant to include compounds that differ only in the presence of isotopically enriched atoms. For example, compounds having the present structures except for the replacement of a hydrogen by a deuterium are expressly included in the present invention.

II. General

[0046] Cathepsin S is a cysteine protease that has been associated with several normal and disease processes in mammals. Specifically, cathepsin S has been directly associated with inflammation, arthritis, and atherosclerosis, as a result of its role in MHC class II antigen presentation. In a preferred aspect, the present invention provides compounds that inhibit the activity of cathepsin S. The present invention also provides methods for treating several disease states in mammals by inhibiting the activity of cathepsin S. In a more preferred aspect, the compounds of the present invention selectively inhibit cathepsin S in the presence of at least one cathepsin isozyme.

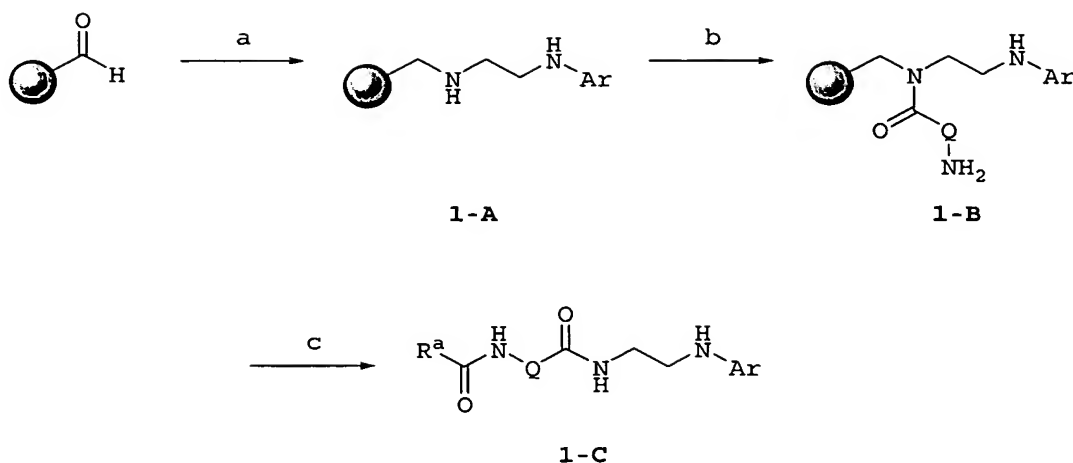
III. Compounds

A. Preparation of Compounds

[0047] In the following schemes, several methods of preparing the compounds of the present invention are illustrative. One of skill in the art will appreciate that these methods are representative, and in no way inclusive of all methods for preparing the compounds of the present invention. The radicals in the schemes are as described in Formula I.

[0048] Compounds of the present invention wherein W is not $R^7-NH-(C=O)-NH-CHR^8$ -, can be made via the route shown in Scheme 1.

Scheme 1

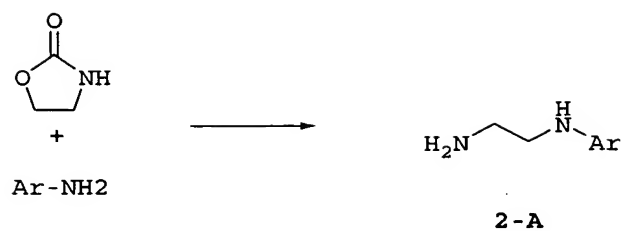


- a) i) $\text{NH}_2\text{CH}_2\text{CH}_2\text{NHAr}$, AcOH, DMF, rt;
 ii) $\text{NaHB}(\text{OAc})_3$, DMF;
 5 b) i) $\text{FmocHNQCO}_2\text{H}$, HOBt, DIC, DMF, rt;
 ii) 20% piperidine in DMF;
 c) i) $\text{R}^a\text{CO}_2\text{H}$, HOBt, DIC, DMF, rt;
 ii) TFA/DCM/ H_2O .

10 **[0049]** As shown therein, polystyrene aldehyde (PAL) resin is reductively aminated with a monoaryl diamine ($\text{NH}_2\text{CH}_2\text{CH}_2\text{NHAr}$) to obtain the resin **1-A** (Scheme 1). This material is acylated with an N-protected amino acid (e.g. $\text{FmocHNQCO}_2\text{H}$) using standard conditions (as described in A. R. Chamberlin, *Chem. Rev.* **1997**, 97, 2243-2266; M. Bodanszky et al. *The Practice of Peptide Synthesis 2nd*, Springer-Verlag, 1984) and the product is then
 15 deprotected with piperidine to furnish **1-B**. After acylation with $\text{R}^a\text{CO}_2\text{H}$ under standard amide coupling condition, cleavage from resin using TFA furnishes **1-C**. In scheme 1, R^a can be R^1 , R^4 , R^6 , R^{10} or R^{11} ; Q can be $-\text{CHR}^2-$, $-\text{CHR}^3-$, $-\text{CHR}^5-$, $-\text{CHR}^9-$, or $-\text{CHR}^{12}-$.

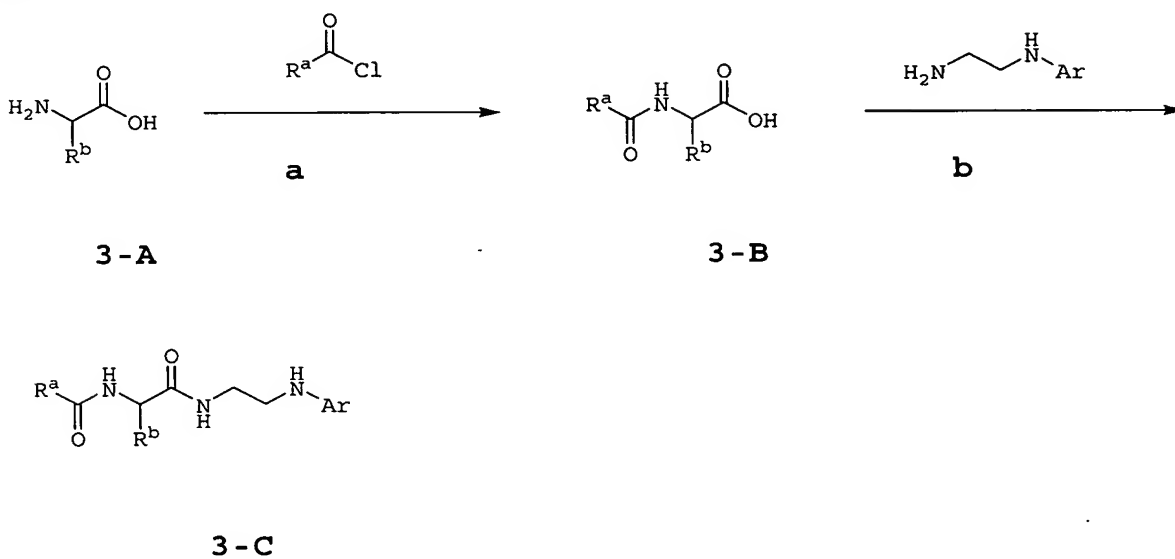
[0050] The arylaminoethylamines **2-A** (Scheme 2) used in the present invention are prepared by a decarboxylative ring opening of oxazolidin-2-one with an aromatic amine as
 20 described in G. S. Poindexter *et al. J. Org. Chem.* **1992**, 57, 6257-65; E. Altman *et al. J. Med Chem.* **2002**, 45, 2352-54 and references cited therein.

Scheme 2



[0051] In an alternate aspect, compounds of the present invention wherein W is not $\text{R}^7\text{-NH-(C=O)-NH-CHR}^8$ can also be prepared according to Scheme 3. In scheme 3, R^a can be R^1 , R^4 , R^6 , R^{10} or R^{11} ; R^b can be R^2 , R^3 , R^5 , R^9 , or R^{12} .

Scheme 3



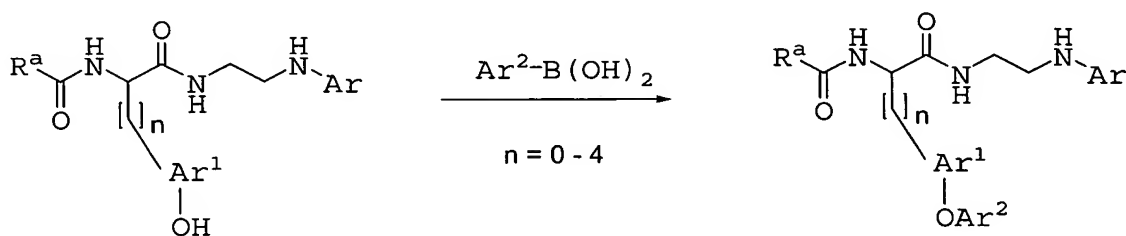
a) NaOH , H_2O ;

b) HOBt , DIC .

[0052] Compounds of the present invention wherein W is $\text{R}^7\text{-NH-(C=O)-NH-CHR}^8$, can be made via the route shown in Scheme 4. In scheme 4, R^a can be R^7 ; Q can be -CHR^8 .

[0054] Compounds of the present invention, wherein R^2 , R^3 , R^5 , R^8 , R^9 or R^{12} consists of, for example, an aryloxy-aryl moiety, can be prepared via copper-promoted arylation of phenols with arylboronic acids (Scheme 6) according to (a) D.A. Evans *et al. Tetrahedron Lett.* **1998**, 39, 2937-40; (b) D. M. Chan *et al. Tetrahedron Lett.* **1998**, 39, 2933-36; and (c) S. V. Ley *et al. Angew. Chem. Int. Ed. Engl.* **2003**, 42, 5400-5449; or Ullman ether synthesis, see (a) S. L. Buchwald *et al. J. Am. Chem. Soc.* **1999**, 121, 4369-4378; (b) J. F. Hartwig *et al. J. Am. Chem. Soc.* **1999**, 121, 3224-3225 and references cited therein. In scheme 6, R^a can be R^1 , R^4 , R^6 , R^{10} or R^{11} .

Scheme 6



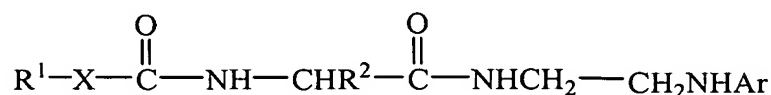
[0055] Synthetic approaches to non-commercially available α - and β -amino acids used in the present invention are widely described in the literature. The typical methods are illustrated in the following references. See: (a) D. J. Ager *et al. Current opinion in drug discovery & development* **2001**, 4, 800-807; (b) R. O. Duthaler *Tetrahedron* **1994**, 50, 1539-1650; (c) M. J. O'Donnell *Aldrichimica Acta* **2001**, 34, 3-15; (d) K. B. Sharpless *et al. J. Am. Chem. Soc.* **1998**, 120, 1207-17; (e) E. Juaristi *et al. Aldrichimica Acta* **1994**, 27, 3-11; (f) D. C. Cole *Tetrahedron* **1994**, 50, 9517-9582, and references cited therein.

[0056] Compounds of the present invention in which W is, for example, R^1 -X-(C=O)-NH-CHR²-, are prepared (a) by Suzuki coupling (N. Miyaura *et al. Synth. Commun.* **1981**, 11(7), 513-19; A. Suzuki *et al. Chem Rev.* **1995**, 95, 2457-2483) of appropriate phenylboronic acid with bromofurancarboxylic acid or bromothiophenecarboxylic acid, when X is furanyl or thienyl; (b) according to Bartoli, J. *et al. J. Med. Chem.* **1998**, 41, 1855-1868 ; Holzer, W. *et al. J. Heterocycl. Chem.* **1993**, 30(4), 865-872 ; Molteni, G. *et al. New J. Chem.* **2002**, 26(10), 1340-1345; Seigo I. *et al. Bioorg. Med. Chem.* **2001**, 11(7), 879-882, when X is pyrazolyl; (c) according to Evans, D. L. *et al. J. Org. Chem.* **1979**, 44, 497; Takeuchi, K. *et al. J. Med. Chem.* **1998**, 41, 5362-5374; Dondoni, A. *et al. J. Org. Chem.* **1987**, 52(15), 3413-3420; Millan, D. S. *et al. Tetrahedron* **2000**, 56(5), 811-816; Duarte, M. P. *et al. Tetrahedron Lett.* **2000**, 41(39), 7433-7435, when X is oxazolyl; and (d) according to Bartoli, J. *et al. J. Med. Chem.* **1998**, 41(11), 1855-1868; Wright, S. W. *et al. J. Med. Chem.* **2002**, 45(18), 3865-

3877; Janusz, J. M. *et al. J. Med. Chem.* **1998**, 41(18), 3515-3529; Tanaka, C. *et al. Chem. Pharm. Bulletin* **1982**, 30(11), 4195-8, when X is thiazolyl.

B. Preferred Compounds

[0057] In one aspect, preferred compounds of the present invention have the following structural formula:



Ia

wherein:

R^1 is a member selected from the group consisting of phenyl substituted with 0-2 R^{1a} , pyridyl substituted with 0-2 R^{1a} , and pyridinium N-oxide substituted with 0-2 R^{1a} ;

each R^{1a} is independently a member selected from the group consisting of Cl, F, OCF_3 , OCH_3 , CH_3 and CF_3 ;

X is a member selected from the group consisting of furanylene substituted with 0-1 R^x , thienylene substituted with 0-1 R^x , pyrazolylene substituted with 0-1 R^x , thiazolylene substituted with 0-1 R^x , and oxazolylene substituted with 0-1 R^x ;

R^x is a member selected from the group consisting of F, Cl, CH_3 and CF_3 ;

R^2 is a member selected from the group consisting of phenyl substituted with 0-2 R^{2a} , and $(CH_2)_nR^{2b}$;

each R^{2a} is independently a member selected from the group consisting of Cl, F, OCF_3 , OCH_3 , CH_3 and CF_3 ;

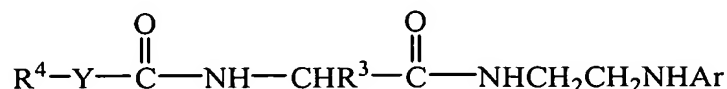
R^{2b} is independently a member selected from the group consisting of phenyl substituted with 0-2 R^{2a} , cyclopentyl, cyclohexyl and tetrahydropyranyl;

n is the integer 1 or 2;

Ar is a phenyl substituted with 0-2 R^{19} ; and

each R^{19} is independently a member selected from the group consisting of F, Cl, $COOH$, C_1 - C_4 alkoxy, $OCHF_2$ and OCF_3 .

[0058] In another aspect, preferred compounds of the present invention have the following structural formula:



Ib

wherein:

R^4 is a member selected from the group consisting of phenyl substituted with 0-3 R^{4a} ,
thienyl, tetrazolyl, cyclopentenyl and indolyl;

5 each R^{4a} is a member selected from the group consisting of phenyl, OH, C_1 - C_4 alkyl,
 C_1 - C_4 alkoxy, CF_3 , OCF_3 , F, Cl, $CH_3S(=O)_2$ -, morpholinyl, pyrrolidinyl,
piperidinyl and 4-acetylpiperazinyl;

Y is a member selected from the group consisting of $-CR^{17}R^{18}$, $-NH-CH_2-$ and
 $-O-CH_2-$;

10 R^3 is $(CH_2)_mR^{3b}$;

R^{3b} is selected from the group consisting of phenyl substituted with 0-2 R^{2a} ,
cyclopentyl and cyclohexyl;

each R^{2a} is independently a member selected from the group consisting of Cl, F,
 OCF_3 , OCH_3 , CH_3 and CF_3 ;

15 m is the integer 1 or 2;

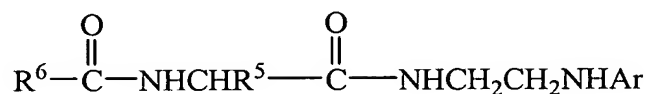
each of R^{17} and R^{18} is independently a member of H, OH, F, phenyl and C_1 - C_3 alkyl;
alternatively, R^{17} and R^{18} may be taken together to form a C_3 - C_6 cycloalkyl;

Ar is a phenyl substituted with 0-2 R^{19} ; and

each R^{19} is independently a member selected from the group consisting of F, Cl,

20 $COOH$, C_1 - C_4 alkoxy, $OCHF_2$ and OCF_3 .

[0059] In still another aspect, preferred compounds of the present invention have the
following structural formula:



Ic

25 wherein:

R^5 is a member selected from the group consisting of phenyl substituted with 0-2 R^{5a} ,
thiophene, naphthyl, and CH_2R^{5b} , CH_2CH_2 (cyclohexyl),
 $CH_2CH_2CH_2$ (cyclohexyl), CH_2CH_2Ph , $CH(CH_3)R^{5c}$, $CH_2CH=CHPh$, -
 CH_2OCH_2Ph , and $-CH(CH_3)OCH_2Ph$;

30 each R^{5a} is independently a member selected from the group consisting of F, Cl, NO_2 ,
 OCH_3 , OCH_2Ph , OPh , CH_3 , OCF_3 and CF_3 ;

R^{5b} is independently a member selected from the group consisting of phenyl substituted with 0-2 R^{5c}; cyclopentyl, cyclohexyl, naphthyl, indolyl and pyridyl;

R^{5c} is independently a member selected from the group consisting of OH, Cl, F, Br, I, CN, NO₂, CH₃, OCH₃, ^tBu, O-^tBu, -NHC(=O)CH₃, CF₃, OCF₃, phenyl substituted with 0-2 R^{5d}, phenoxy substituted with 0-2 R^{5d}, benzyloxy substituted with 0-2 R^{5d}, pyridyl substituted with 0-2 R^{5d}, pyrimidinyl substituted with 0-2 R^{5d}, and thienyl substituted with 0-2 R^{5d};

R^{5d} is independently a member selected from the group consisting of CH₃, Cl, F, OCH₃, CF₃, OCF₃, N(CH₃)₂, acetyl, OH, CH₂OH, NH₂, CN and NO₂;

R^{5e} is phenyl substituted with 0-2 R^{5a};

R⁶ is a member selected from the group consisting of phenyl substituted with 0-3 R^{6a}, furanyl substituted with 0-2 R^{6b}; thienyl substituted with 0-2 R^{6b}; oxazolyl substituted with 0-2 R^{6b}; thiazolyl substituted with 0-2 R^{6b}; pyridyl, pyridazinyl and cyclopropyl;

each R^{6a} is independently a member selected from the group consisting of Cl, F, Br, OCF₃, CF₃, C₁-C₄ alkyl, C₁-C₄ alkoxy, -S(=O)₂CH₃, CN, -N(CH₃)₂, OCF₂H, -CH₂-benzimidazole, -NH-S(=O)₂CH₃, -NR¹³R¹⁴, OR¹⁴, CH₂-morpholine, CH₂NH₂, OCH₂Ph, and OPh;

alternatively, two R^{6a} substituents on adjacent atoms may be combined to form a 5 to 6 membered heterocyclic fused radical, wherein said 5 to 6 membered heterocyclic fused radical has 1 or 2 oxygen atom(s);

each R^{6b} is independently a member selected from the group consisting of NH₂, F, Cl, Br, -S(=O)₂R¹⁵, CH₃, and CF₃;

R¹³ is a member selected from the group consisting of H and C₁-C₄ alkyl;

R¹⁴ is pyrimidinyl substituted with 0-2 R¹⁶;

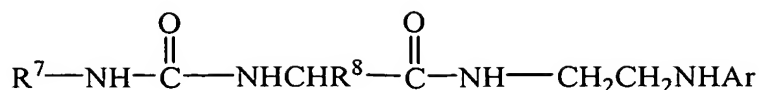
R¹⁵ is a member selected from the group consisting of C₁-C₄ alkyl, morpholinyl, pyrrolidinyl and piperidinyl;

R¹⁶ is a member selected from the group consisting of CH₃ and OCH₃;

Ar is a phenyl substituted with 0-2 R¹⁹; and

each R¹⁹ is independently a member selected from the group consisting of F, Cl, COOH, C₁-C₄ alkoxy, OCHF₂ and OCF₃.

[0060] In still yet another aspect, preferred compounds of the present invention have the following structural formula:



Id

wherein:

R^7 is a member selected from the group consisting of $(CH_2)_p R^{7a}$; and naphthyl substituted with 0-2 R^{7b} ;

p is the integer 0, 1, or 2;

R^{7a} is a member selected from the group consisting of phenyl substituted with 0-2 R^{7b} ;

R^{7b} is a member selected from the group consisting of F, Cl, CF_3 , C_1 - C_4 alkyl, C_1 - C_4 alkoxy, OCF_3 , phenoxy and acetyl;

alternatively, two R^{7b} substituents on adjacent atoms may be combined to form a 5 to 6 membered heterocyclic fused radical, wherein said 5 to 6 membered heterocyclic fused radical has 1 or 2 oxygen atom(s);

R^8 is $-CH_2-R^{3b}$;

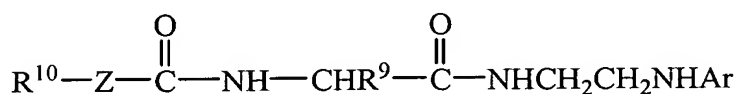
R^{3b} is selected from the group consisting of phenyl substituted with 0-2 R^{2a} , cyclopentyl and cyclohexyl;

each R^{2a} is independently a member selected from the group consisting of Cl, F, OCF_3 , OCH_3 , CH_3 and CF_3 ;

Ar is a phenyl substituted with 0-2 R^{19} ; and

each R^{19} is independently a member selected from the group consisting of F, Cl, $COOH$, C_1 - C_4 alkoxy, $OCHF_2$ and OCF_3 .

[0061] In another aspect, preferred compounds of the present invention have the following structural formula:



Ie

wherein:

R^{10} is a member selected from the group consisting of phenyl substituted with 0-2 R^{10a} , 5 membered heteroaryl containing 1 to 4 heteroatoms each independently a member selected from the group consisting of N, O and S, wherein said heteroaryl is substituted with 0-2 R^{10a} , 6 membered heteroaryl containing 1 to

2 N, wherein said heteroaryl is substituted with 0-2 R^{10a}, morpholinyl substituted with 0-2 R^{10a}, piperazinyl substituted with 0-2 R^{10a} and piperidinyl substituted with 0-2 R^{10a};

each R^{10a} is independently a member selected from the group consisting of Cl, F, C₁-

C₄ alkyl, C₁-C₄ alkoxy, OCF₃, and CF₃;

alternatively, two R^{10a} substituents on adjacent atoms may be combined to form a 5 to 6 membered heterocyclic fused radical, wherein said 5 to 6 membered heterocyclic fused radical comprises 1 or 2 heteroatom(s);

Z is phenylene;

R⁹ is (CH₂)_qR^{9a};

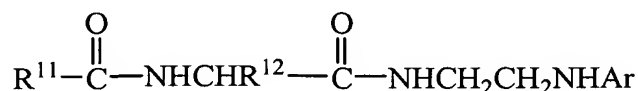
R^{9a} is a member selected from the group consisting of cyclopentyl, phenyl and cyclohexyl;

q is the integer 1 or 2;

Ar is a phenyl substituted with 0-2 R¹⁹; and

each R¹⁹ is independently a member selected from the group consisting of F, Cl, COOH, C₁-C₄ alkoxy, OCHF₂ and OCF₃.

[0062] In still another aspect, preferred compounds of the present invention have the following structural formula:



wherein:

R¹¹ is a member selected from the group consisting of indolyl substituted with 0-2 R^{11a}; benzofuranyl substituted with 0-2 R^{11a}; benzothienyl substituted with 0-2 R^{11a}; benzothiazole substituted with 0-2 R^{11a}; benzisoxazolyl substituted with 0-2 R^{11a}; benzoxazolyl substituted with 0-2 R^{11a}; and pyrazolo[1,5-a]pyrimidinyl substituted with 0-2 R^{11a}; piperidinyl N-substituted with 0-1 R^{11b}; morpholinyl N-substituted with 0-1 R^{11b}; and 2-oxo-pyrrolidinyl with 0-1 R^{11b};

each R^{11a} is independently a member selected from the group consisting of Cl, F, NH₂, CH₃, OCH₃, -C(=O)OCH₃, OCF₃, and CF₃;

each R^{11b} is independently a member selected from the group consisting of
pyrimidinyl substituted with 0-2 R^{11c}; benzyl, acetyl, CH₂-furanlyl, and CH₂-
thienyl;

each R^{11c} is independently a member selected from the group consisting of Br and
5 CH₃;

R¹² is (CH₂)_sR^{12a};

R^{12a} is a member selected from the group consisting of cyclopentyl and cyclohexyl;
s is the integer 1 or 2;

Ar is a phenyl substituted with 0-2 R¹⁹; and

10 each R¹⁹ is independently a member selected from the group consisting of F, Cl,
COOH, C₁-C₄ alkoxy, OCHF₂ and OCF₃.

[0063] Preferred compounds of Formula I are set forth below in Table I:

TABLE I

1. N-((S)-1-(2-(4-methoxyphenylamino)ethylcarbamoyl)-3-phenylpropyl)-5-(3-
15 (trifluoromethyl)phenyl)furan-2-carboxamide;
2. N-((S)-1-(2-(4-methoxyphenylamino)ethylcarbamoyl)-2-(2-chlorophenyl)ethyl)-5-(3-
(trifluoromethyl)phenyl)furan-2-carboxamide;
3. N-((S)-1-(2-(4-methoxyphenylamino)ethylcarbamoyl)-2-(3-chlorophenyl)ethyl)-5-(3-
(trifluoromethyl)phenyl)furan-2-carboxamide;
- 20 4. N-((S)-1-(2-(4-methoxyphenylamino)ethylcarbamoyl)-2-(4-chlorophenyl)ethyl)-5-(3-
(trifluoromethyl)phenyl)furan-2-carboxamide;
5. N-((S)-1-(2-(4-methoxyphenylamino)ethylcarbamoyl)-2-(tetrahydro-2H-pyran-4-
yl)ethyl)-5-(3-(trifluoromethyl)phenyl)furan-2-carboxamide;
6. N-((S)-1-(2-(4-methoxyphenylamino)ethylcarbamoyl)-2-cyclopentylethyl)-5-(3-
25 (trifluoromethyl)phenyl)furan-2-carboxamide;
7. (S)-N-{2-[4-(2,3-Dimethyl-phenoxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-
ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
8. (±)-N-((2-(4-methoxyphenylamino)ethylcarbamoyl)(4-chlorophenyl)methyl)-3-
methylbenzamide;

9. (±)-N-((2-(4-methoxyphenylamino)ethylcarbamoyl)(phenyl)-methyl)-5-(3-(trifluoromethyl)phenyl)furan-2-carboxamide;
10. N-((S)-1-(2-(4-(difluoromethoxy)phenylamino)ethylcarbamoyl)-2-cyclohexylethyl)-5-(3-(trifluoromethyl)phenyl)furan-2-carboxamide;
- 5 11. 4-[2-(3-Cyclohexyl-2-(S)-{[5-(3-trifluoromethyl-phenyl)-furan-2-carbonyl]-amino}-propionylamino)-ethylamino]-benzoic acid;
12. 2-[2-(3-Cyclohexyl-2-(S)-{[5-(3-trifluoromethyl-phenyl)-furan-2-carbonyl]-amino}-propionylamino)-ethylamino]-benzoic acid;
- 10 13. 4-Cyclohexyl-2-(S)-(2-(R)-phenyl-propionylamino)-N-[2-(4-trifluoromethoxy-phenylamino)-ethyl]-butyramide;
14. Acetyl-piperidine-4-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
15. (S)-2-{2-[4-(4-Acetyl-piperazin-1-yl)-phenoxy]-acetylamino}-3-cyclohexyl-N-[2-(4-trifluoromethoxy-phenylamino)-ethyl]-propionamide;
- 15 16. (S)-2-Chloro-N-{1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-3-methyl-benzamide;
17. Cyclohexyl-2-[2-(4-methoxy-phenyl)-acetylamino]-N-[2-(4-trifluoromethoxy-phenylamino)-ethyl]-propionamide;
18. (S)-N-{2-[4-(3,5-Dichloro-phenoxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
- 20 19. N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-methanesulfonyl-benzamide;
20. (S)-4-Benzoyloxy-N-{1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-3,5-dimethyl-benzamide;
- 25 21. (S)-4-Methoxy-N-{1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-3,5-dimethyl-benzamide;
22. 5-Methoxy-1H-indole-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;

23. (S)-3-Bromo-N-{1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-4-methyl-benzamide;
24. Furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 5 25. Thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
26. Furan-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 10 27. N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-(5-methyl-[1,2,4]oxadiazol-3-yl)-benzamide;
28. 5-(4-Fluoro-phenyl)-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
29. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-2,4,5-trimethyl-benzamide;
- 15 30. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-2,4,5-trimethyl-benzamide;
31. 5-(3-Fluoro-phenyl)-furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
32. Benzyl-morpholine-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 20 33. (S)-N-{2-[4-(4-Dimethylamino-phenoxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
34. 2'-Chloro-biphenyl-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 25 35. 5-(2-Trifluoromethyl-phenyl)-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
36. 5-(3-Fluoro-phenyl)-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;

37. Thiophene-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
38. Oxo-1-thiophen-2-ylmethyl-pyrrolidine-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 5 39. Furan-2-ylmethyl-5-oxo-pyrrolidine-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
40. Methyl-5-(pyrrolidine-1-sulfonyl)-furan-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 10 41. (S)-1-Phenyl-5-trifluoromethyl-1H-pyrazole-4-carboxylic acid {1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-amide;
42. 5-p-Tolyl-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
43. Benzoimidazol-1-ylmethyl-N-{2-cyclohexyl-1-(S)-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide;
- 15 44. (S)-1-(4-Chloro-phenyl)-5-trifluoromethyl-1H-pyrazole-4-carboxylic acid {1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-amide;
45. (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-p-tolyloxy-phenyl)-ethyl]-3-methyl-benzamide;
- 20 46. Cyclohexyl-2-(S)-(2-tetrazol-1-yl-acetylamino)-N-[2-(4-trifluoromethoxy-phenylamino)-ethyl]-propionamide;
47. 5-m-Tolyl-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
48. 2,7-Dimethyl-pyrazolo[1,5-a]pyrimidine-6-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 25 49. 2-Methyl-5-(morpholine-4-sulfonyl)-furan-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
50. 5-(3-Trifluoromethyl-phenyl)-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;

51. 5-m-Tolyl-furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
52. (S)-2,3-Dihydro-benzofuran-7-carboxylic acid {1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-amide;
- 5 53. Methanesulfonyl-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
54. 2-Phenyl-thiazole-4-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
55. (S)-3-Cyano-N-{1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-benzamide;
- 10 56. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-3-(2-methyl-thiazol-4-yl)-benzamide;
57. (S)-N-[2-(4-Methoxy-phenylamino)-ethyl]-3-phenyl-2-(3-phenyl-ureido)-propionamide;
- 15 58. 3-Cyclohexyl-2-(S)-(2-(S)-hydroxy-2-phenyl-acetyl-amino)-N-[2-(4-trifluoromethoxy-phenylamino)-ethyl]-propionamide;
59. Benzo[c]isoxazole-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
60. N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-difluoromethoxy-benzamide;
- 20 61. N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-isopropoxy-benzamide;
62. Phenyl-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 25 63. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-nicotinamide;
64. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-isonicotinamide;

65. Phenyl-furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
66. (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-o-tolyloxy-phenyl)-ethyl]-3-methyl-benzamide;
- 5 67. N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-oxazol-5-yl-benzamide;
68. 5-(3-Trifluoromethyl-phenyl)-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
69. 5-(2-Trifluoromethyl-phenyl)-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 10 70. 5-p-Tolyl-furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
71. N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-[(4,6-dimethyl-pyrimidin-2-yl)-methyl-amino]-benzamide;
- 15 72. 1-(4,6-Dimethyl-pyrimidin-2-yl)-piperidine-4-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
73. N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-(4,6-dimethoxy-pyrimidin-2-yloxy)-benzamide;
74. 3'-Methoxy-biphenyl-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 20 75. N-{3-Cyclohexyl-1-(S)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-2-(R)-phenyl-butyramide;
76. 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-(2-thiophen-2-yl-acetylamino)-propionamide;
- 25 77. 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-(2-thiophen-3-yl-acetylamino)-propionamide;
78. (S)-3-Bromo-N-{1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-benzamide;

79. Acetyl-piperidine-4-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
80. N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-(4,6-dimethoxy-pyrimidin-2-yl)-benzamide;
- 5 81. 1-(5-Bromo-pyrimidin-2-yl)-piperidine-4-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
82. (S)-2-(2-Cyclopent-2-enyl-acetyl-amino)-N-[2-(4-methoxy-phenylamino)-ethyl]-3-phenyl-propionamide;
83. Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(2-1H-indol-3-yl-acetyl-amino)-propionamide;
- 10 84. N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-3-methanesulfonylamino-benzamide;
85. 5-(3-Trifluoromethyl-phenyl)-furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 15 86. Cyclohexyl-2-(S)-(2-(R,S)-fluoro-2-phenyl-acetyl-amino)-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide;
87. Cyclohexyl-N-(S)-[2-(4-fluoro-phenylamino)-ethyl]-2-[2-(4-trifluoromethoxy-phenyl)-acetyl-amino]-propionamide;
88. (S)-2-[3-(4-Chloro-phenyl)-ureido]-N-[2-(4-methoxy-phenylamino)-ethyl]-3-phenyl-propionamide;
- 20 89. (S)-N-[2-(4-Methoxy-phenylamino)-ethyl]-2-[3-(4-phenoxy-phenyl)-ureido]-3-phenyl-propionamide;
90. (S)-N-[2-(4-Methoxy-phenylamino)-ethyl]-2-(3-phenethyl-ureido)-3-phenyl-propionamide;
- 25 91. (S)-2-[3-(4-Fluoro-benzyl)-ureido]-N-[2-(4-methoxy-phenylamino)-ethyl]-3-phenyl-propionamide;
92. N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-(4,6-dimethyl-pyrimidin-2-ylamino)-benzamide;

93. 1-(5-Bromo-pyrimidin-2-yl)-piperidine-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
94. (S)-2-(3-Benzo[1,3] dioxol-5-yl-ureido)-N-[2-(4-methoxy-phenylamino)-ethyl]-3-phenyl-propionamide;
- 5 95. 3-Cyclohexyl-2-(S)-[2-(2,5-difluorophenyl)-acetylamino]-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide;
96. (S)-2-[3-(3-Fluoro-benzyl)-ureido]-N-[2-(4-methoxy-phenylamino)-ethyl]-3-phenyl-propionamide;
97. (S)-N-[2-(4-Methoxy-phenylamino)-ethyl]-3-phenyl-2-(3-o-tolyl-ureido)-propionamide;
- 10 98. (S)-N-{2-[4-(3,4-Dichloro-phenoxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
99. 3-Cyclohexyl-2-(S)-[2-(3,4-difluoro-phenyl)-acetylamino]-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide;
- 15 100. 3-Cyclohexyl-2-(S)-[2-(2,4-difluoro-phenyl)-acetylamino]-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide;
101. (S)-N-[2-(4-Methoxy-phenylamino)-ethyl]-2-(3-naphthalen-1-yl-ureido)-3-phenyl-propionamide;
102. (S)-2-[3-(2-tert-Butyl-6-methyl-phenyl)-ureido]-N-[2-(4-methoxy-phenylamino)-ethyl]-3-phenyl-propionamide;
- 20 103. (S)-2-[3-(4-Acetyl-phenyl)-ureido]-N-[2-(4-methoxy-phenylamino)-ethyl]-3-phenyl-propionamide;
104. (S)-N-[2-(4-Methoxy-phenylamino)-ethyl]-2-[3-(3-methoxy-phenyl)-ureido]-3-phenyl-propionamide;
- 25 105. (S)-Biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
106. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-trifluoromethyl-benzamide;

107. 2-(S)-[2-(2-Chloro-4-fluoro-phenyl)-acetylamino]-3-cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide;
108. (S)-2-Chloro-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
- 5 109. (S)-4-Benzoyloxy-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide;
110. (S)-4-Benzoyloxy-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3,5-dimethyl-benzamide;
111. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-methoxy-3,5-dimethyl-benzamide;
- 10 112. (S)-3-Bromo-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-methyl-benzamide;
113. (S)-5-Fluoro-1H-indole-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 15 114. (S)-2-Amino-4-methyl-thiazole-5-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
115. (S)-1-Phenyl-5-trifluoromethyl-1H-pyrazole-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
116. (S)-1-(4-Chloro-phenyl)-5-trifluoromethyl-1H-pyrazole-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 20 117. (S)-5-(3-Trifluoromethyl-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
118. (S)-3-Chloro-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide;
- 25 119. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-dimethylamino-benzamide;
120. (S)-3-Cyano-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide;

121. (S)-4-Cyano-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide;
122. N-{2-cyclohexyl-1-(S)-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-(R)-phenyl-propionamide;
- 5 123. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-(2-methyl-thiazol-4-yl)-benzamide;
124. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-[1,2,4]triazol-1-yl-benzamide;
125. 3-Cyclohexyl-2-(S)-[2-(3,5-difluoro-phenyl)-acetylamino]-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide;
- 10 126. (S)-N-{3-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-3-trifluoromethyl-benzamide;
127. (S)-N-{3-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-4-morpholin-4-yl-benzamide;
- 15 128. (4-Cyclohexyl-N-[2-(4-methoxy-phenylamino)-ethyl]-2-(S)-(2-(S)-phenyl-propionylamino)-butyramide;
129. (S)-4-Benzoyloxy-N-{3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-benzamide;
130. (S)-Biphenyl-4-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide;
- 20 131. (S)-5-Chloro-1H-indole-2-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide;
132. (S)-5-Fluoro-1H-indole-2-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide;
- 25 133. (S)-2-Amino-4-methyl-thiazole-5-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide;
134. (S)-5-Chloro-benzofuran-2-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide;

135. N-{2-cyclohexyl-1-(S)-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-(R)-phenyl-butyramide;
136. (S)-5-(3-Trifluoromethyl-phenyl)-furan-2-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide;
- 5 137. (S)-Benzothiazole-6-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide;
138. (S)-N-{3-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-3-trifluoromethoxy-benzamide;
139. (S)-3-Cyano-N-{3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-benzamide;
- 10 140. (S)-4-Cyano-N-{3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-benzamide;
141. N-{2-cyclohexyl-1-(S)-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-phenoxy-benzamide;
- 15 142. (S)-N-{3-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-3-(2-methyl-thiazol-4-yl)-benzamide;
143. (S)-N-{3-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-4-[1,2,4]triazol-1-yl-benzamide;
144. (S)-Biphenyl-3-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide;
- 20 145. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-fluoro-benzamide;
146. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3,4-difluoro-benzamide;
- 25 147. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-fluoro-2-methyl-benzamide;
148. (S)-2-Chloro-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-5-methyl-benzamide;

149. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-fluoro-3-trifluoromethyl-benzamide;
150. (S)-5-Methyl-1-phenyl-1H-pyrazole-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 5 151. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-propyl-benzamide;
152. 3-Cyclohexyl-2-(S)-[2-(4-fluoro-phenyl)-acetylamino]-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide;
153. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-methoxy-benzamide;
- 10 154. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-fluoro-5-trifluoromethyl-benzamide;
155. (S)-3-Chloro-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-fluoro-benzamide;
- 15 156. (S)-5-Chloro-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-fluoro-benzamide;
157. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-5-fluoro-2-methyl-benzamide;
158. (S)-1-Phenyl-cyclopropanecarboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 20 159. (S)-3-Cyclohexyl-N-[2-(4-methoxy-phenylamino)-ethyl]-2-(2-phenylamino-acetylamino)-propionamide;
160. 3-Cyclohexyl-2-(S)-(2-(R)-hydroxy-2-phenyl-acetylamino)-N-[2-(4-methoxy-phenylamino)-ethyl]-propionamide ;
- 25 161. (S)-1-(4-Fluoro-phenyl)-5-methyl-1H-pyrazole-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
162. (S)-1-(4-Methoxy-phenyl)-5-methyl-1H-pyrazole-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;

163. (S)-1-(4-Chloro-phenyl)-5-methyl-1H-pyrazole-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
164. N-{2-Cyclohexyl-1-(S)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-(S)-phenyl-butyramide;
- 5 165. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-fluoro-5-trifluoromethyl-benzamide;
166. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-fluoro-3-trifluoromethyl-benzamide;
167. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-fluoro-3-methyl-benzamide;
- 10 168. (S)-5-(4-Chloro-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
169. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-fluoro-4-trifluoromethyl-benzamide;
- 15 170. (S)-4'-Chloro-biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
171. (S)-3', 5'-Dichloro-biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
172. (S)-3'-Methoxy-biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 20 173. (S)-3'-Chloro-biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
174. (S)-2'-Methoxy-biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 25 175. (S)-4'-Chloro-biphenyl-3-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
176. (S)-4-Benzo[1,3]dioxol-5-yl-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide;

177. (S)-5-Bromo-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
178. (S)-3,5-Dibromo-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide;
- 5 179. (S)-3,5-Dichloro-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide;
180. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3,5-dimethoxy-benzamide;
- 10 181. (S)-Biphenyl-3-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
182. (S)-5-Bromo-thiophene-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
183. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-phenoxy-benzamide;
- 15 184. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-phenoxy-benzamide;
185. (S)-1H-Indole-3-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
186. (S)-Benzothiazole-6-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 20 187. (S)-2-Amino-benzothiazole-6-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
188. (S)-4-Methyl-2-(4-trifluoromethyl-phenyl)-thiazole-5-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 25 189. (S)-4-(4-Chloro-phenyl)-thiophene-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
190. (S)-2-Methyl-5-trifluoromethyl-oxazole-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;

191. (S)-4-(4-Methoxy-phenyl)-thiophene-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
192. N-{2-Cyclohexyl-1-(S)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-(S)-phenyl-propionamide;
- 5 193. (S)-5-(2-Chloro-5-trifluoromethyl-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
194. (S)-2'-Chloro-biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
195. (S)-1-(5-Bromo-pyrimidin-2-yl)-piperidine-4-carboxylic acid {2-cyclohexyl-10 1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
196. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-(4,6-dimethyl-pyrimidin-2-ylamino)-benzamide;
197. (S)-1-(5-Bromo-pyrimidin-2-yl)-piperidine-3-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 15 198. (S)-3'-Fluoro-biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
199. (S)-3-Aminomethyl-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide;
200. (S)-N-{2-Cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-morpholin-4-ylmethyl-benzamide;
- 20 201. (S)-5-(2-Fluoro-phenyl)-thiophene-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
202. (S)-N-{3-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-3-methyl-benzamide;
- 25 203. (S)-N-{4-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-butyl}-3-methyl-benzamide;
204. (S)-N-{[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-phenyl-methyl}-3-methyl-benzamide;

205. (S)-5-(4-Trifluoromethyl-phenyl)-thiophene-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
206. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-3-phenyl-propyl}-3-methyl-benzamide;
- 5 207. (S)-5-(4-Trifluoromethoxy-phenyl)-thiophene-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
208. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-4-phenyl-but-3-enyl}-3-methyl-benzamide;
- 10 209. (S)-5-(3-Trifluoromethoxy-phenyl)-thiophene-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
210. (S)-5-(2-Methoxy-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
211. N-{(4-Methoxy-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide;
- 15 212. (S)-5-(2-Fluoro-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
213. (S)-5-(4-Trifluoromethyl-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 20 214. N-{(2-Benzoyloxy-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide;
215. (S)-5-(4-Trifluoromethoxy-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
216. N-{(2-Chloro-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide;
- 25 217. N-{(4-Benzoyloxy-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide;
218. N-{[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-naphthalen-1-yl-methyl}-3-methyl-benzamide;

219. (S)-5-(2-Trifluoromethyl-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
220. N- {[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-o-tolyl-methyl}-3-methyl-benzamide;
- 5 221. (S)-N- {2-Cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-[1,2,4]triazol-1-yl-benzamide;
222. N- {(2,4-Dichloro-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide;
- 10 223. N- {(2,3-Dichloro-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide;
224. N- {(2,4-Dimethyl-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide;
225. N- {(2,4-Dimethoxy-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide;
- 15 226. N- {[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-thiophen-2-yl-methyl}-3-methyl-benzamide;
227. N- {(4-Fluoro-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide;
- 20 228. (S)-N- {2-(4-Hydroxy-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
229. (S)-N- {2-(2,4-Dichloro-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
230. (S)-N- {2-(3,5-Difluoro-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
- 25 231. (S)-N- {2-(3,4-Dichloro-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
232. (S)-4-Benzyloxy-N- {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide;

233. (S)-N-{2-(4-Acetylamino-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
234. (S)-Biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 5 235. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-p-tolyl-ethyl}-3-methyl-benzamide;
236. (S)-N-{2-(3-Fluoro-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
- 10 237. (S)-N-{2-(3,4-Difluoro-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
238. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-m-tolyl-ethyl}-3-methyl-benzamide;
239. (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(2-trifluoromethyl-phenyl)-ethyl]-3-methyl-benzamide;
- 15 240. (S)-N-{2-(4-Cyano-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
241. (S)-N-{2-(4-Bromo-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
242. (S)-N-{2-(4-Iodo-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
- 20 243. (S)-N-{2-(4-Chloro-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
244. (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-nitro-phenyl)-ethyl]-3-methyl-benzamide;
- 25 245. (S)-N-{2-(4-Fluoro-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
246. (S)-5-(3-Trifluoromethyl-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;

247. (S)-N-{2-(4-Benzoyloxy-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
248. (S)-N-{2-[4-(2,6-Dichloro-benzyloxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
- 5 249. (S)-N-{2-(4-Methoxy-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
250. 2-Amino-4-methyl-thiazole-5-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
251. (S)-5-(2-Chloro-5-trifluoromethyl-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 10 252. (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(3-trifluoromethyl-phenyl)-ethyl]-3-methyl-benzamide;
253. (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-trifluoromethyl-phenyl)-ethyl]-3-methyl-benzamide;
- 15 254. (S)-N-{2-Benzoyloxy-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
255. (S)-N-{2-(4-tert-Butyl-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
256. Cyclohexyl-N-[2-(4-methoxy-phenylamino)-ethyl]-2-(S)-(2-(S)-phenyl-propionylamino)-butyramide;
- 20 257. (S)-N-{2-(1H-Indol-3-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
258. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-naphthalen-1-yl-ethyl}-3-methyl-benzamide;
- 25 259. (S)-N-{2-Benzoyloxy-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-3-methyl-benzamide;
260. 3-Cyclohexyl-2-(S)-[2-(3-fluoro-phenyl)-acetylamino]-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide;

261. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-naphthalen-2-yl-ethyl}-3-methyl-benzamide;
262. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-pyridin-3-yl-ethyl}-3-methyl-benzamide;
- 5 263. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-pyridin-4-yl-ethyl}-3-methyl-benzamide;
264. Furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 10 265. 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-(2-tetrazol-1-yl-acetyl-amino)-propionamide;
266. N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-nitro-phenyl)-propyl]-3-methyl-benzamide;
267. (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-m-tolyloxy-phenyl)-ethyl]-3-methyl-benzamide;
- 15 268. threo-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-propyl}-3-methyl-benzamide;
269. erythro-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-propyl}-3-methyl-benzamide;
- 20 270. (S)-N-{2-Biphenyl-4-yl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
271. (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(3'-nitro-biphenyl-4-yl)-ethyl]-3-methyl-benzamide;
272. Furan-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 25 273. (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(2'-nitro-biphenyl-4-yl)-ethyl]-3-methyl-benzamide;
274. (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-pyridin-3-yl-phenyl)-ethyl]-3-methyl-benzamide;

275. (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-thiophen-3-yl-phenyl)-ethyl]-3-methyl-benzamide;
276. (S)-N-{2-(4'-Cyano-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
- 5 277. (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-pyridin-4-yl-phenyl)-ethyl]-3-methyl-benzamide;
278. (S)-N-{2-(4'-Chloro-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
279. (S)-N-{2-(2',3'-Dimethoxy-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
- 10 280. (S)-N-{2-(3'-Amino-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
281. (S)-N-{2-(3',4'-Dimethoxy-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
- 15 282. (S)-N-{2-(4'-Hydroxymethyl-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
283. (S)-N-{2-(5'-Fluoro-2'-methoxy-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
284. (S)-N-{2-(3'-Hydroxymethyl-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
- 20 285. (S)-N-{2-(2',5'-Dimethoxy-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
286. (S)-N-{2-(2',5'-Dichloro-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
- 25 287. (S)-N-{2-(4'-Dimethylamino-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
288. (S)-N-{2-(2'-Acetyl-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;

289. (S)-N-{2-(4'-Hydroxy-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
290. (S)-N-{2-(3'-Acetyl-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
- 5 291. (S)-N-{2-[4-(2,4-Dimethoxy-pyrimidin-5-yl)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
292. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-[4-(6-methoxy-pyridin-3-yl)-phenyl]-ethyl}-3-methyl-benzamide;
- 10 293. Methanesulfonyl-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
294. N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-2-(S)-phenyl-propionamide;
295. Pyridazine-4-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 15 296. N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-3-methanesulfonyl-benzamide;
297. 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-(2-1H-tetrazol-5-yl-acetylamino)-propionamide;
- 20 298. Cyclopropanecarboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
299. N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-4-methanesulfonylamino-benzamide;
300. (S)-N-{2-[4-(4-Chloro-phenoxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
- 25 301. 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-[2-(4-methoxy-phenyl)-acetylamino]-propionamide;
302. 2-(S)-[2-(3-Chloro-phenyl)-acetylamino]-3-cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide;

303. 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-phenylacetylamino-propionamide;
304. 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-(2-o-tolyl-acetylamino)-propionamide;
- 5 305. 2-(S)-[2-(4-Chloro-phenyl)-acetylamino]-3-cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide;
306. 3-Cyclohexyl-2-(S)-[2-(2-fluoro-phenyl)-acetylamino]-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide;
- 10 307. 3-Cyclohexyl-2-(S)-diphenylacetylamino-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide;
308. N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-2-(2-fluoro-biphenyl-4-yl)-propionamide;
309. N-{2-cyclohexyl-1-(S)-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-2-p-tolyl-propionamide;
- 15 310. N-{2-cyclohexyl-1-(S)-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-2-(4-fluoro-phenyl)-propionamide;
311. N-{2-cyclohexyl-1-(S)-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-2-(4-hydroxy-phenyl)-propionamide;
- 20 312. 2-(4-Chloro-phenyl)-N-{2-cyclohexyl-1-(S)-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-propionamide;
313. N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-4-methanesulfonyl-benzamide;
314. Thiazole-4-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
- 25 315. N-{2-cyclohexyl-1-(S)-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-2-(R)-phenyl-propionamide;
316. 4-Cyano-N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide;

317. 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-(2-(R)-hydroxy-2-phenyl-acetylamino)-propionamide;
318. N-{2-cyclohexyl-1-(S)-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-2-(R)-phenyl-butyramide;
- 5 319. Phenyl-cyclopropanecarboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
320. N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-2-(R,S)-(4-fluoro-phenyl)-propionamide;
- 10 321. Cyano-N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide;
322. 5-(4-Fluoro-phenyl)-furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide;
323. Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-[2-(3-trifluoromethyl-phenyl)-acetylamino]-propionamide;
- 15 324. Cyano-N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide;
325. 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-[2-(4-trifluoromethyl-phenyl)-acetylamino]-propionamide;
- 20 326. 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-[2-(4-methanesulfonyl-phenyl)-acetylamino]-propionamide;
327. (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-phenoxy-phenyl)-ethyl]-3-methyl-benzamide;
328. (S)-N-{2-[4-(4-Methoxy-phenoxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
- 25 329. (S)-N-{2-[4-(3-Chloro-phenoxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide;
330. (S)-N-{2-[4-(3,5-Dimethyl-phenoxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide.

[0064] Compounds of the present invention are either obtained in the free form, or as a salt thereof if salt forming groups are present, or as esters if ester forming groups are present.

[0065] Compounds of the present invention that have acidic groups can be converted into salts with pharmaceutically acceptable bases, e.g., an aqueous alkali metal hydroxide, advantageously in the presence of an ethereal or alcoholic solvent, such as a lower alkanol. Resulting salts can be converted into the free compounds, e.g., by treatment with acids.

These, or other salts can also be used for purification of the compounds obtained.

Ammonium salts are obtained by reaction with the appropriate amine, e.g., diethylamine, and the like.

[0066] In certain aspects, compounds of the present invention having basic groups can be converted into acid addition salts, especially pharmaceutically acceptable salts. These are formed, for example, with inorganic acids, such as mineral acids, for example, sulfuric acid, a phosphoric or hydrohalic acid, or with organic carboxylic acids, such as (C₁-C₄) alkane carboxylic acids which, for example, are unsubstituted or substituted by halogen, for example, acetic acid, such as saturated or unsaturated dicarboxylic acids, for example, oxalic, succinic, maleic or fumaric acid, such as hydroxycarboxylic acids, for example, glycolic, lactic, malic, tartaric or citric acid, such as amino acids, for example, aspartic or glutamic acid, or with organic sulfonic acids, such as (C₁-C₄)-alkylsulfonic acids (for example, methanesulfonic acid) or arylsulfonic acids which are unsubstituted or substituted (for example, by halogen). Preferred are salts formed with hydrochloric acid, methanesulfonic acid and maleic acid.

[0067] In view of the close relationship between the free compounds and the compounds in the form of their salts or esters, whenever a compound is referred to in this context, a corresponding salt or ester is also intended, provided such is possible or appropriate under the circumstances.

[0068] The compounds, including their salts, can also be obtained in the form of their hydrates, or include other solvents used for their crystallization.

[0069] The compounds of the present invention that comprise free hydroxyl groups may also exist in the form of pharmaceutically acceptable, physiologically cleavable esters, and as such are included within the scope of the invention. Such pharmaceutically acceptable esters are preferably prodrug ester derivatives, such being convertible by solvolysis or cleavage under physiological conditions to the corresponding compounds of the present invention

which comprise free hydroxyl groups. Suitable pharmaceutically acceptable prodrug esters are those derived from a carboxylic acid, a carbonic acid monoester or a carbamic acid, preferably esters derived from an optionally substituted lower alkanolic acid or an arylcarboxylic acid.

5 [0070] As will be apparent to one of skill in the art, certain compounds of the present invention possess asymmetric carbon atoms (optical centers) or double bonds; the racemates, diastereomers, enantiomers, geometric isomers and individual isomers are all intended to be encompassed within the scope of the present invention.

10 [0071] The present invention provides compounds which inhibit cathepsin S selectively. In certain preferred aspects, the present invention provides compounds which selectively inhibit cathepsin S in the presence of cathepsin isozymes, such as cathepsin A, B, C, D, E, F, G, H, K, L, M, O, P, Q, R, V, W, X and combinations thereof. In a more preferred aspect, the present invention provides compounds which selectively inhibit cathepsin S in the presence of cathepsin K.

15 [0072] Compounds of the present invention useful for treating cathepsin S dependent conditions, preferably have cathepsin S inhibition constants less than 10 μM . More preferably, compounds of the present invention useful for treating cathepsin S dependent conditions have cathepsin S inhibition constants of less than 1.0 μM . Most preferably, compounds of the present invention useful for treating cathepsin S dependent conditions have
20 cathepsin S inhibition constants of less than 0.1 μM .

[0073] In a preferred aspect, compounds of the present invention that selectively inhibit cathepsin S in the presence of a cathepsin isozyme (*e.g.* cathepsin K), have a cathepsin isozyme inhibition constant at least 10 times greater than their cathepsin S inhibition constant. In a more preferred aspect, compounds of the present invention that selectively
25 inhibit cathepsin S in the presence of a cathepsin isozyme, have a cathepsin isozyme inhibition constant at least 100 times greater than their cathepsin S inhibition constant. In a most preferred aspect, compounds of the present invention that selectively inhibit cathepsin S in the presence of a cathepsin isozyme, have a cathepsin isozyme inhibition constant at least 1000 times greater than their cathepsin S inhibition constant.

IV. Compositions

[0074] The pharmaceutical compositions according to the invention are those suitable for enteral, such as oral or rectal, transdermal, topical, and parenteral administration to mammals, including humans, to inhibit cathepsin S activity, and for the treatment of cathepsin S dependent disorders, in particular chronic neuropathic pain (see, WO 03/020287), Alzheimer's disease and certain autoimmune disorders, including, but not limited to, juvenile onset diabetes, multiple sclerosis, pemphigus vulgaris, Graves' disease, myasthenia gravis, systemic lupus erythematosus, rheumatoid arthritis and Hashimoto's thyroiditis; allergic disorders, including, but not limited to, asthma; and allogeneic immune responses, including, but not limited to, rejection of organ transplants or tissue grafts.

[0075] More particularly, the pharmaceutical compositions comprise an effective cathepsin S inhibiting amount of a compound of the present invention.

[0076] The pharmacologically active compounds of the present invention are useful in the manufacture of pharmaceutical compositions comprising an effective amount thereof in conjunction or mixture with excipients or carriers suitable for either enteral or parenteral application.

[0077] Preferred are tablets and gelatin capsules comprising the active ingredient together with a) diluents, e.g., lactose, dextrose, sucrose, mannitol, sorbitol, cellulose and/or glycine; b) lubricants, e.g., silica, talcum, stearic acid, its magnesium or calcium salt and/or polyethyleneglycol; for tablets also c) binders, e.g., magnesium aluminum silicate, starch paste, gelatin, tragacanth, methylcellulose, sodium carboxymethylcellulose and or polyvinylpyrrolidone; if desired d) disintegrants, e.g., starches, agar, alginic acid or its sodium salt, or effervescent mixtures; and/or e) absorbents, colorants, flavors and sweeteners. Injectable compositions are preferably aqueous isotonic solutions or suspensions, and suppositories are preferably prepared from fatty emulsions or suspensions. The compositions may be sterilized and/or contain adjuvants, such as preserving, stabilizing, wetting or emulsifying agents, solution promoters, salts for regulating the osmotic pressure and/or buffers. In addition, they may also contain other therapeutically valuable substances. The compositions are prepared according to conventional mixing, granulating or coating methods, respectively, and contain about 0.1 to 75%, preferably about 1 to 50%, of the active ingredient.

[0078] Tablets may be either film coated or enteric coated according to methods known in the art.

[0079] Suitable formulations for transdermal application include an effective amount of a compound of the present invention with carrier. Preferred carriers include absorbable pharmacologically acceptable solvents to assist passage through the skin of the host. For example, transdermal devices are in the form of a bandage comprising a backing member, a reservoir containing the compound optionally with carriers, optionally a rate controlling barrier to deliver the compound to the skin of the host at a controlled and predetermined rate over a prolonged period of time, and means to secure the device to the skin. Matrix transdermal formulations may also be used.

[0080] Suitable formulations for topical application, e.g., to the skin and eyes, are preferably aqueous solutions, ointments, creams or gels well-known in the art. Such may contain solubilizers, stabilizers, tonicity enhancing agents, buffers and preservatives.

[0081] The pharmaceutical formulations contain an effective cathepsin S inhibiting amount of a compound of the present invention as defined above, either alone or in combination with another therapeutic agent.

[0082] In conjunction with another active ingredient, a compound of the present invention may be administered either simultaneously, before or after the other active ingredient, either separately by the same or different route of administration or together in the same pharmaceutical formulation.

[0083] The dosage of active compound administered is dependent on the species of warm-blooded animal (mammal), the body weight, age and individual condition, and on the form of administration. A unit dosage for oral administration to a mammal of about 50 to 70 kg may contain between about 5 and 500 mg of the active ingredient.

[0084] In a preferred aspect, the pharmaceutical composition of the present invention provides a compound according to Formula I.

[0085] In one aspect of the present invention, compositions of the present invention that comprise compounds of the present invention and pharmaceutically acceptable excipients, selectively inhibit cathepsin S in the presence of other cathepsin isozymes. In a more preferred aspect, the present invention provides compositions which selectively inhibit cathepsin S in the presence of cathepsin K.

[0086] In another aspect of the present invention, compositions of the present invention useful for treating cathepsin S dependent conditions, preferably have cathepsin S inhibition constants less than 10 μ M. More preferably, compositions of the present invention useful for treating cathepsin S dependent conditions have cathepsin S inhibition constants of less than 1.0 μ M. Most preferably, compositions of the present invention useful for treating cathepsin S dependent conditions have cathepsin S inhibition constants of less than 0.1 μ M.

[0087] In a preferred aspect, compositions of the present invention utilize compounds that selectively inhibit cathepsin S in the presence of a cathepsin isozyme (*e.g.* cathepsin K), have a cathepsin isozyme inhibition constant at least 10 times greater than their cathepsin S inhibition constant. In a more preferred aspect, compounds of the present invention that selectively inhibit cathepsin S in the presence of cathepsin isozyme, have a cathepsin isozyme inhibition constant at least 100 times greater than their cathepsin S inhibition constant. In a most preferred aspect, compounds of the present invention that selectively inhibit cathepsin S in the presence of cathepsin isozyme, have a cathepsin isozyme inhibition constant at least 1000 times greater than their cathepsin S inhibition constant.

V. Methods

[0088] In view of their activity as inhibitors of cathepsin S, compounds of the present invention are particularly useful in mammals as agents for treatment and prophylaxis of diseases and medical conditions involving elevated levels of cathepsin S. For example, the compounds of the present invention are useful in treating Alzheimer's disease and certain autoimmune disorders, including, but not limited to juvenile onset diabetes, multiple sclerosis, pemphigus vulgaris, Graves' disease, myasthenia gravis, systemic lupus erythematosus, rheumatoid arthritis and Hashimoto's thyroiditis; allergic disorders, including, but not limited to asthma; and allogeneic immune responses, including, but not limited to, rejection of organ transplants or tissue grafts.

[0089] Beneficial effects are evaluated *in vitro* and *in vivo* pharmacological tests generally known in the art, and as illustrated herein.

[0090] The above cited properties are demonstrable *in vitro* and *in vivo* tests, using advantageously mammals, *e.g.*, rats, mice, dogs, rabbits, monkeys or isolated organs and tissues, as well as mammalian enzyme preparations, either natural or prepared by, *e.g.*, recombinant technology. Compounds of the present invention can be applied *in vitro* in the

form of solutions, e.g., preferably aqueous solutions or suspensions, and *in vivo* either enterally or parenterally, preferably orally, e.g., as a suspension or in aqueous solution, or as a solid capsule formulation. The dosage *in vitro* may range between about 10^{-5} molar and 10^{-9} molar concentrations. The dosage *in vivo* may range, depending on the route of administration, between about 0.1 and 100 mg/kg.

[0091] The antiarthritic efficacy of the compounds of the present invention for the treatment of rheumatoid arthritis can be determined using models such as, or similar to, the rat model of adjuvant arthritis, as described previously (R. E. Esser, *et al.*, *J. Rheumatology* **1993**, *20*, 1176). The efficacy of the compounds of the present invention for the treatment of osteoarthritis can be determined using models such as, or similar to, the rabbit partial lateral meniscectomy model, as described previously (Colombo *et al.*, *Arth. Rheum.* **1993**, *26*, 875-886). The efficacy of the compounds in the model can be quantified using histological scoring methods, as described previously (O'Byrne *et al.*, *Inflamm. Res.* **1995**, *44*, S 177-S118).

[0092] The present invention also relates to methods of using compounds of the present invention and their pharmaceutically acceptable salts, or pharmaceutical compositions thereof, in mammals for inhibiting cathepsin S, and for the treatment of cathepsin S dependent conditions, such as the cathepsin S dependent conditions described herein, e.g., inflammation, rheumatoid arthritis and osteoarthritis.

[0093] In a preferred aspect, the present invention relates to a method of treating rheumatoid arthritis, osteoarthritis, and inflammation (and other diseases as identified above) in mammals comprising administering to a mammal in need thereof, a correspondingly effective amount of a compound of the present invention.

[0094] In a preferred aspect, the method of the present invention provides a compound according to Formula I.

[0095] Methods of the present invention useful for treating cathepsin S dependent conditions, preferably use compounds that have cathepsin S inhibition constants less than 10 μ M. More preferably, methods of the present invention useful for treating cathepsin S dependent conditions use compounds that have cathepsin S inhibition constants of less than 1.0 μ M. Most preferably, methods of the present invention useful for treating cathepsin S

dependent conditions use compounds that have cathepsin S inhibition constants of less than 0.1 μ M.

[0096] Moreover, the present invention relates to a method of selectively inhibiting cathepsin S activity in a mammal which comprises administering to a mammal in need thereof, an effective cathepsin S inhibiting amount of a compound of the present invention. In a preferred aspect, the methods of the present invention use compounds that selectively inhibit cathepsin S in the presence of a cathepsin isozyme, such as cathepsin A, B, C, D, E, F, G, H, K, L, M, O, P, Q, R, V, W and X. In a more preferred aspect, methods of the present invention use compounds that selectively inhibit cathepsin S in the presence of cathepsin K.

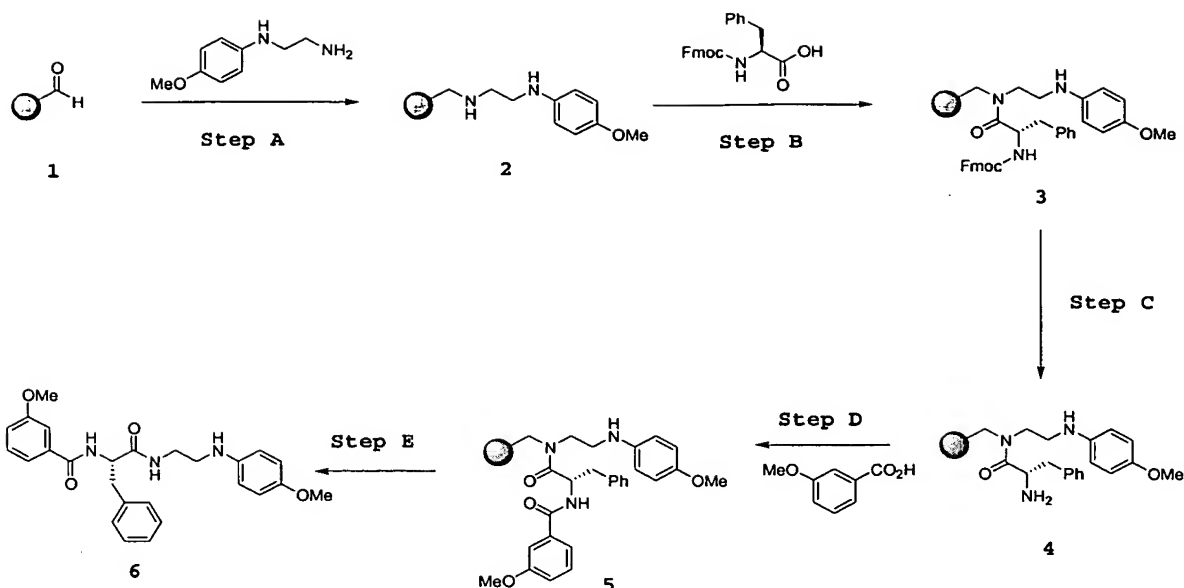
[0097] In a preferred aspect, methods of the present invention use compounds that selectively inhibit cathepsin S in the presence of a cathepsin isozyme (*e.g.* cathepsin K), have a cathepsin isozyme inhibition constant at least 10 times greater than their cathepsin S inhibition constant. In a more preferred aspect, compounds of the present invention that selectively inhibit cathepsin S in the presence of cathepsin isozyme, have a cathepsin isozyme inhibition constant at least 100 times greater than their cathepsin S inhibition constant. In a most preferred aspect, compounds of the present invention that selectively inhibit cathepsin S in the presence of cathepsin isozyme, have a cathepsin isozyme inhibition constant at least 1000 times greater than their cathepsin S inhibition constant.

VI. Examples

A. Compounds

[0098] **General Procedure.** All solvents stated as anhydrous were purchased that way from the manufacturer and used as received. All other purchased reagents were used as received. Unless otherwise stated, all reactions were carried out under a positive pressure of nitrogen. Silica gel chromatography was performed using pre-packed cartridges and an instrument for making a linear solvent gradient along with automated fraction collection. ^1H NMR spectral data were reported as follows: chemical shift on the δ scale (using residual protio solvent as the internal standard), multiplicity (s = singlet, d = doublet, t = triplet, q = quartet, m = multiplet), integration and coupling constant in hertz. ^{13}C spectra were recorded as APT experiments and were reported in ppm with residual solvent for internal standard.

[0099] **Preparation 1.** Solid phase synthesis (S)-3-methoxy-N-{1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-benzamide.



Step A: An aldehyde-functionalized polystyrene resin 1 (“Pal-Resin”, 16.76 g @1.05 mmol/g, 17.6mmol) was swelled in DMF (50 mL) for 10 min. N1-(4-methoxy-phenyl)-ethane-1,2-diamine (5.85g, 35 mmol) in DMF (150 mL) was added followed by acetic acid (8.1 mL, 8 eq), and the mixture was agitated for 1 h at room temperature. Sodium triacetoxyborohydride (11.2g, 52.8 mmol eq.) was then added and the mixture was shaken for 16 hours at room temperature. The reductively aminated resin 2 was then filtered and washed (DMF \times 3, equal mixture of methanol/dichloromethane \times 4, Acetonitrile \times 3).

Step B: The resin 2 (17.6 mmol) was swelled in DMF (50 mL) and a solution of Fmoc-Phe-OH (20.45 g, 3 eq), HOBt (8.08 g, 3 eq) and DIC (4.58 mL, 3eq) was added. The mixture was shaken for 3 h, then washed (DMF \times 3, equal mixture of methanol/dichloromethane \times 4, Acetonitrile \times 3).

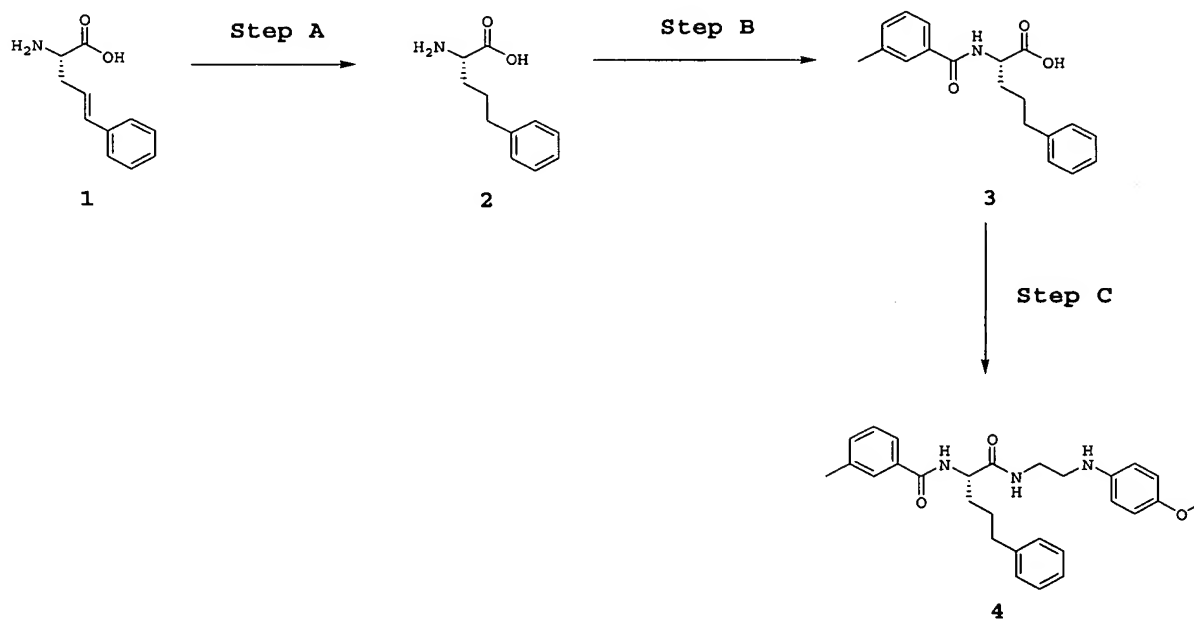
Step C: The resin 3 (117 mg, 0.077 mmol) was weighed into reaction vessels followed by a stirrer bar then treated with piperidine in DMF (4 mL of a 20% solution) and the mixture agitated for 1 h. The resin is then washed (3 \times DMF, 3 \times dichloromethane) to yield resin 4.

Step D: Resin 4 (0.077 mmol) was swelled in DMF (1 mL) and a premixed solution of hydroxybenzotriazole (35 mg, 0.23 mmol), 3-methoxybenzoic acid (35 mg, 0.23 mmol) and diisopropylcarbodiimide (20 μ L, 0.231 mmol) in DMF (3.0 mL) is added. The reaction is agitated for three hours and then washed (DMF \times 4, then dichloromethane \times 3) and dried under nitrogen to yield resin 5.

Step E: The resin **5** (0.077 mmol) was treated with a mixture of trifluoroacetic acid, dichloromethane and water (45:45:10, v/v, 4 mL). The resin was agitated for one hour then retreated, agitated again for 5 min, then washed with the above cleavage mixture and filtered into vials. The solvent was evaporated *in vacuo* and the title compound purified using a

5 Waters mass-directed reverse phase LCMS system (flow 7.5 min method, gradient 10-90% acetonitrile/water with 0.35% trifluoroacetic acid). Solvent was removed by lyophilization to afford the title compound as a white solid. (9 mg, 0.02mmol, 26%). ¹H NMR (CD₃OD) δ(ppm) 7.38(d, *J*= 9.0Hz, 2H), 7.31(m, 2H), 7.28(m, 5H), 7.21(m, 1H), 7.10(m, 1H), 7.05(d, *J*=9.0Hz, 2H), 4.64(dd, *J*=7.1Hz, *J*=8.4Hz, 1H), 3.82(s, 3H), 3.81(s, 3H), 3.60(m, 1H),
10 3.44(m, 1H), 3.37(m, 2H), 3.22(dd, *J*=8.6Hz, *J*=13.6Hz, 1H), 3.12(dd, *J*=7.0Hz, *J*=13.6Hz, 1H); C₂₆H₂₉N₃O₄; LCMS: 448.2 (M+H)⁺.

[0100] Preparation 2. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-4-phenyl-butyl}-3-methyl-benzamide.



15 **Step A:** L-Styryl alanine **1** (50 mg, 0.26 mmol) was dissolved in MeOH (5 mL) and the reaction vessel was flushed with nitrogen. Catalytic amount of palladium (10% on carbon) was added and the reaction vessel was placed under a hydrogen atmosphere. The mixture was stirred for 2 h at room temperature, then filtered over celite. The organic solvent was removed in vacuo to yield (S)-2-amino-5-phenyl-pentanoic acid **2** (51 mg, quant.) as a white
20 powder: ¹H-NMR (400MHz, CD₃OD) δ = 7.42-7.25 (m, 5H), 3.71 (t, *J*= 6.0, 1H), 2.69 (t,

$J = 6.4$, 2H), 1.85 (m, 2H), 1.70 (m, 2H). MS calcd. for $C_{11}H_{16}NO_2$ ($M+H^+$) 194.12, found 194.4.

Step B: (S)-2-Amino-5-phenyl-pentanoic acid **2** (23 mg, 0.12 mmol) was dissolved in H_2O (1 mL) containing equimolar amounts of NaOH (5 mg, 0.12 mmol). The solution was cooled to 0 °C, then m-Toluic acid chloride (16 μL , 0.12 mmol) was added dropwise under vigorous stirring. The mixture was allowed to warm to room temperature and stirred for approx. 12 h. After acidification with 1 M HCl (1 mL), the product was extracted from the reaction mixture with DCM (4 mL). The organic layer was separated and the solvent was removed in vacuo to yield (S)-2-(3-methyl-benzoylamino)-5-phenyl-pentanoic acid **3** (21 mg, 56%) as a white solid: 1H -NMR (400MHz, CD_3OD) δ = 7.83-7.09 (m, 9H), 4.62 (dd, $J = 5.0$, $J = 9.1$, 1H), 2.65 (m, 2H), 2.37 (s, 3H), 2.01-1.67 (m, 4H). MS calcd. for $C_{19}H_{22}NO_3$ ($M+H^+$) 312.16, found 312.4.

Step C: (S)-2-(3-Methyl-benzoylamino)-5-phenyl-pentanoic acid **3** (21 mg, 0.07 mmol) was dissolved in DCM (2 mL), HOBT (20 mg, 0.14 mmol) and DIC (23 μL , 0.14 mmol) were added and the solution was stirred for 10 min at room temperature. N-(4-Methoxyphenyl)-ethane-1,2-diamine (24 mg, 0.14 mmol) was added and the solution was stirred for 3 h at room temperature. The solvent was removed in vacuo, and the remainder was purified by reverse HPLC to afford the title compound (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-4-phenyl-butyl}-3-methyl-benzamide **4** (18 mg, 0.04 mmol, 56%) as a white solid: 1H -NMR (400MHz, CD_3OD) δ = 7.56-7.48 (m, 2H), 7.36-7.16 (m, 7H), 6.96-6.87 (m, 4H), 4.48 (dd, $J = 6.2$, $J = 13.7$, 1H), 3.82 (s, 3H), 3.78-3.64 (m, 2H), 3.58-3.50 (m, 2H), 2.71-2.64 (m, 2H), 2.35 (s, 3H), 2.06-1.70 (m, 4H). MS calcd. for $C_{28}H_{34}N_3O_3$ ($M+H^+$) 460.26, found 460.5.

* Compounds prepared according to Preparation 1.

\$ Compounds prepared according to Example 57.

Compounds prepared according to Example 10.

** Compounds prepared according to Example 227.

Compounds prepared according to Example 270.

\$\$ Compounds prepared according to Example 327.

[0101] **Example 1.** N-((S)-1-(2-(4-methoxyphenylamino)ethylcarbamoyl)-3-phenylpropyl)-5-(3-(trifluoromethyl)phenyl)furan-2-carboxamide. * C₃₁H₃₀F₃N₃O₄; LCMS: 566.5 (M+H)⁺.

[0102] **Example 2.** N-((S)-1-(2-(4-methoxyphenylamino)ethylcarbamoyl)-2-(2-chlorophenyl)ethyl)-5-(3-(trifluoromethyl)phenyl)furan-2-carboxamide. * C₃₀H₂₇ClF₃N₃O₄; LCMS: 586.4 (M+H)⁺.

[0103] **Example 3.** N-((S)-1-(2-(4-methoxyphenylamino)ethylcarbamoyl)-2-(3-chlorophenyl)ethyl)-5-(3-(trifluoromethyl)phenyl)furan-2-carboxamide. * C₃₀H₂₇ClF₃N₃O₄; LCMS: 586.4 (M+H)⁺.

[0104] **Example 4.** N-((S)-1-(2-(4-methoxyphenylamino)ethylcarbamoyl)-2-(4-chlorophenyl)ethyl)-5-(3-(trifluoromethyl)phenyl)furan-2-carboxamide. * C₃₀H₂₇ClF₃N₃O₄; LCMS: 586.4 (M+H)⁺.

[0105] **Example 5.** N-((S)-1-(2-(4-methoxyphenylamino)ethylcarbamoyl)-2-(tetrahydro-2H-pyran-4-yl)ethyl)-5-(3-(trifluoromethyl)phenyl)furan-2-carboxamide. * C₂₉H₃₂F₃N₃O₅; LCMS: 560.4 (M+H)⁺.

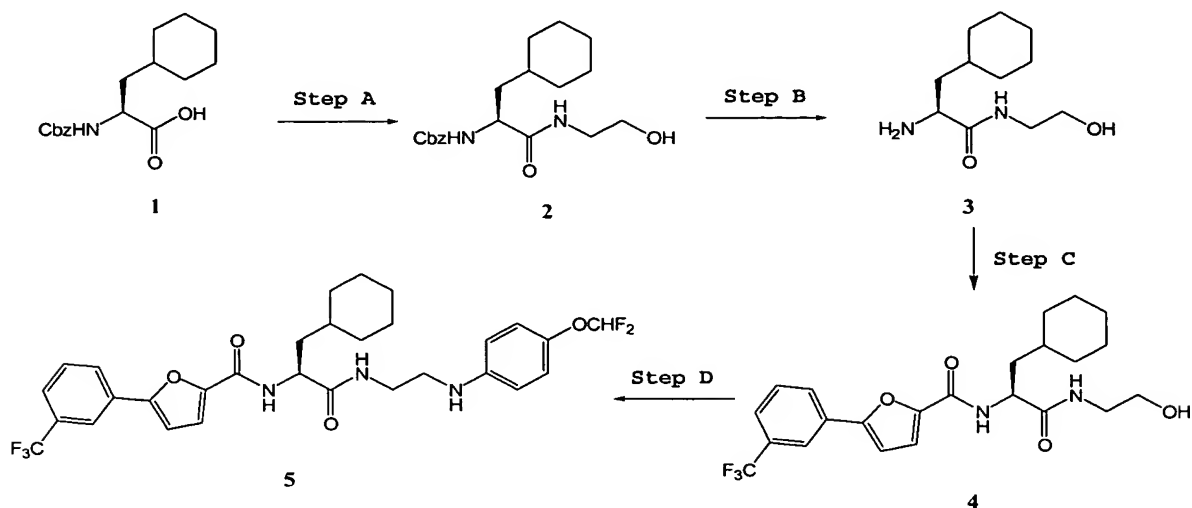
[0106] **Example 6.** N-((S)-1-(2-(4-methoxyphenylamino)ethylcarbamoyl)-2-cyclopentylethyl)-5-(3-(trifluoromethyl)phenyl)furan-2-carboxamide. * C₂₉H₃₂F₃N₃O₄; LCMS: 544.5 (M+H)⁺.

[0107] **Example 7.** (S)-N-{2-[4-(2,3-Dimethyl-phenoxy)-phenyl]-1-[2-(4-methoxyphenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. ^{ss} Following the procedure of Example 327, except substituting phenyl-boronic acid for 2,3-dimethylphenyl-boronic acid in Step D, the title compound was prepared as a white solid (5 mg, 18%): ¹HNMR (400MHz, CD₃OD) δ = 7.59-6.64 (m, 15H), 4.62 (dd, J = 7.2, J = 8.3, 1H), 3.81 (s, 3H), 3.61-3.04 (m, 6H), 2.37 (s, 3H), 2.29 (s, 3H), 2.05 (s, 3H). MS calcd. for C₃₄H₃₈N₃O₄ (M+H)⁺ 552.29, found 552.4.

[0108] **Example 8.** (±)-N-((2-(4-methoxyphenylamino)ethylcarbamoyl)(4-chlorophenyl)methyl)-3-methylbenzamide. ** C₂₅H₂₆ClN₃O₃; LCMS: 452.4 (M+H)⁺.

[0109] **Example 9.** (±)-N-((2-(4-methoxyphenylamino)ethylcarbamoyl)(phenyl)-methyl)-5-(3-(trifluoromethyl)phenyl)furan-2-carboxamide. ** C₂₉H₂₆F₃N₃O₄; LCMS: 538.4 (M+H)⁺.

[0110] Example 10. *N*-((*S*)-1-(2-(4-(difluoromethoxy)phenylamino)ethylcarbamoyl)-2-cyclohexylethyl)-5-(3-(trifluoromethyl)phenyl)furan-2-carboxamide.



Step A: (*S*)-2-Benzoyloxycarbonylamino-3-cyclohexyl-propionic acid **1** (2.065 g, 6.77 mmol, 1.0 eq.) was dissolved in CH₂Cl₂ (50 mL). *N*-(3-Dimethylaminopropyl)-*N'*-ethylcarbodiimide hydrochloride (EDC, 1.56 g, 8.11 mmol, 1.2 eq.) and 1-Hydroxybenzotriazole hydrate (HOBT, 1.10 g, 8.16 mmol, 1.2 eq.) were added to the reaction slurry. After 20 minutes, ethanolamine (1.6 mL, 26.58 mmol, 3.9 eq.) was added via syringe and the reaction was allowed to stir at room temperature and monitored by LC/MS. After the reaction was allowed to proceed to completion, the reaction mixture was extracted with water (40 mL), 1M HCl (30 mL), followed by saturated NaHCO₃ and saturated NaCl. The organic layer was dried over MgSO₄ and filtered. The organic solvent was removed in vacuo and purified by silica gel chromatography to yield (*S*)-[2-Cyclohexyl-1-(2-hydroxy-ethylcarbamoyl)-ethyl]-carbamic acid benzyl ester **2** (780 mg, 2.24 mmol, 28%) as a white solid: ¹H-NMR (400MHz, CDCl₃) δ 0.76-0.92 (m, 2H), 1.02-1.35 (m, 4H), 1.44-1.73 (m, 7H), 3.17-3.25 (m, 1H), 3.34-3.40 (m, 1H), 3.51-3.69 (m, 3H), 4.24 (dt, 1H, *J* = 8.8 Hz, 5.6 Hz), 4.94 (d, 1H, *J* = 12.0 Hz), 5.07 (d, 1H, *J* = 12.0 Hz), 5.93 (1H, *J* = 12.0 Hz), 7.24-7.30 (m, 5H).

Step B: (*S*)-[2-Cyclohexyl-1-(2-hydroxy-ethylcarbamoyl)-ethyl]-carbamic acid benzyl ester **2** (630 mg, 1.81 mmol) was dissolved in dry MeOH (15 mL) and Pd/C (15 mg, catalytic) was added under N₂. The nitrogen atmosphere was replaced with a H₂ balloon and allowed to stir at room temperature. The reaction was judged to be complete by LC/MS and the reaction was purged with nitrogen and filtered through celite. The solvent was removed in vacuo to yield (*S*)-2-Amino-3-cyclohexyl-N-(2-hydroxy-ethyl)-propionamide **3** (380 mg, 1.77 mmol, 98%) as a clear oil: MS calcd. for C₁₁H₂₂N₂O₂ (M + H⁺) 215.2, found 215.4.

Step C: (*S*)-2-Amino-3-cyclohexyl-N-(2-hydroxy-ethyl)-propionamide **3** (216 mg, 1.0 mmol, 1.0 eq.), *O*-(7-Azabenzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate (HATU, 364 mg, 1.11 mmol, 1.1 eq.), and 5-(3-Trifluoromethyl-phenyl)-furan-2-carboxylic acid (284 mg, 1.11 mmol, 1.1 eq.) was dissolved in CH₂Cl₂ (10 mL, 0.1M) and the solution was stirred for 10 min at room temperature. *N,N*-Diisopropylethylamine (DIPEA, 500 μ L, 2.88 mmol, 2.9 eq.) was added and the solution was stirred at room temperature until all starting materials were consumed by LC/MS. The solvent was removed in vacuo and the residue purified by silica gel chromatography (0% to 40% EtOAc gradient in hexanes) to provide (*S*)-5-(3-Trifluoromethyl-phenyl)-furan-2-carboxylic acid [2-cyclohexyl-1-(2-hydroxy-ethylcarbamoyl)-ethyl]-amide **4** (293 mg, 0.65 mmol, 64%) as a white solid: MS calcd. for C₂₃H₂₇F₃N₂O₄ (M + H⁺) 453.2, found 453.4.

Step D: (*S*)-5-(3-Trifluoromethyl-phenyl)-furan-2-carboxylic acid [2-cyclohexyl-1-(2-hydroxy-ethylcarbamoyl)-ethyl]-amide **4** (170 mg, 0.38 mmol) was dissolved in CH₂Cl₂ (5 mL, 0.07M) under a N₂ atmosphere. Dess-Martin Periodinane (210 mg, 0.49 mmol, 1.3 eq.) was added in one portion and allowed the reaction to stir at room temperature for 3 hours. After the reaction was judged to be complete by TLC, the reaction was diluted with EtOAc (50 mL) and extracted with 1M sodium thiosulfate (30 mL). The organic layer was extracted with saturated NaHCO₃ and saturated NaCl. The organic layer was dried over MgSO₄ and filtered. The organic solvent was removed in vacuo and the resulting aldehyde (158 mg, 0.35 mmol, 93%) was used directly without storage: R_f = 0.67 (1:1 hexanes:EtOAc). The aldehyde (43 mg, 0.10 mmol) was dissolved in MeOH (2.5 mL, 0.04M) and brought to 0°C in an ice bath. 4-Difluoromethoxyaniline (50 μ L, 0.31 mol, 3.3 eq.) and acetic acid (20 mL, 0.34 mmol, 3.6 eq.) were added via syringe followed by sodium cyanoborohydride (20 mg, 0.32 mmol) in one portion. The clear reaction mixture was allowed to slowly warm to room temperature and monitored to completion by LC/MS. The reaction was worked up by rotary evaporation of MeOH, dilution with EtOAc (20 mL) and water (20 mL). The organic phase was separated and washed with 1M NaOH (15 mL) and saturated NaCl (15 mL). The organic layer was dried over MgSO₄, filtered, and concentrated by rotary evaporation. Purification by mass-directed HPLC, evaporation and lyophilization provided (*S*)-5-(3-Trifluoromethyl-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-difluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide **5** as a white amorphous solid (15 mg, 0.02 mmol, 22%): ¹H NMR (CD₃OD, 400 MHz) δ 0.95-1.06 (m, 2H), 1.19-1.29 (m, 3H), 1.40-1.44 (m, 1H), 1.66-1.86 (m, 7H), 3.26-3.29 (m, 2H), 3.43-3.46 (m, 2H), 4.64 (dd, 1H, *J* =

9.2, 6.0 Hz), 6.59 (t, 1H, $J = 74.8$ Hz), 6.76 (d, 2H, $J = 8.8$ Hz), 6.95 (d, 2H, $J = 8.8$ Hz), 7.11 (s, 1H), 7.29 (d, 1H, $J = 3.6$ Hz), 7.66 (s, 2H), 8.13 (s, 1H), 8.25 (s, 1H); HPLC-MS calcd. for $C_{30}H_{32}F_3N_3O_4$ ($M + H^+$) 594.2, found 594.5.

[0111] Example 11. 4-[2- (3-Cyclohexyl-2-(*S*)-{[5-(3-trifluoromethyl-phenyl)-furan-2-carbonyl]-amino}-propionylamino)-ethylamino]-benzoic acid.[#]

¹H NMR (CD_3OD , 400 MHz) δ 0.95-1.06 (m, 2H), 1.19-1.29 (m, 3H), 1.40-1.44 (m, 1H), 1.63-1.85 (m, 7H), 3.31-3.35 (m, 2H), 3.39-3.48 (m, 2H), 4.64 (dd, 1H, $J = 9.6, 5.6$ Hz), 6.62 (d, 2H, $J = 8.8$ Hz), 7.10 (d, 1H, $J = 8.0$ Hz), 7.28 (d, 1H, $J = 7.2$ Hz), 7.62-7.66 (m, 2H), 7.75 (d, 2H, $J = 8.8$ Hz), 8.12-8.14 (m, 1H), 8.29 (s, 1H); HPLC-MS calcd. for $C_{30}H_{32}F_3N_3O_5$ ($M + H^+$) 572.2, found 572.5.

[0112] Example 12. 2-[2- (3-Cyclohexyl-2-(*S*)-{[5-(3-trifluoromethyl-phenyl)-furan-2-carbonyl]-amino}-propionylamino)-ethylamino]-benzoic acid.[#]

¹H NMR (CD_3OD , 400 MHz) δ 0.93-1.02 (m, 2H), 1.17-1.29 (m, 3H), 1.32-1.48 (m, 1H), 1.64-1.86 (m, 7H), 3.31-3.44 (m, 3H), 3.49-3.53 (m, 1H), 4.68 (t, 1H, $J = 7.6$ Hz), 6.57 (t, 1H, $J = 15.2$ Hz), 6.83 (d, 1H, $J = 8.4$ Hz), 7.09 (d, 1H, $J = 3.6$ Hz), 7.27 (d, 1H, $J = 3.6$ Hz), 7.32-7.36 (m, 1H), 7.61-7.65 (m, 2H), 7.86 (dd, 1H, $J = 8.0, 1.6$ Hz), 8.11-8.13 (m, 1H), 8.23 (s, 1H); HPLC-MS calcd. for $C_{30}H_{32}F_3N_3O_5$ ($M + H^+$) 572.2, found 572.5.

[0113] Example 13. 4-Cyclohexyl-2-(*S*)-(2-(*R*)-phenyl-propionylamino)-N-[2-(4-trifluoromethoxy-phenylamino)-ethyl]-butyramide; $C_{28}H_{36}F_3N_3O_3$; ¹H NMR ($CDCl_3$) δ (ppm) 6.97(m, 5H), 6.74(m, 2H), 6.42(m, 1H), 6.27(m, 2H), 5.85(d, $J = 8.4$ Hz, 1H), 3.95(m, 1H), 3.32(m, 1H), 3.11(m, 2H), 2.86(m, 2H), 1.52(m, 1H), 1.35(m, 5H), 1.21(m, 4H), 0.83(m, 6H), 0.49(m, 2H); LCMS: 520.6 ($M + H$)⁺.

[0114] Example 14. 1-Acetyl-piperidine-4-carboxylic acid (*S*)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide. * $C_{26}H_{37}F_3N_4O_4$; ¹H NMR ($CDCl_3$) δ (ppm) 7.29(m, 1H), 7.08(d, $J = 8.4$ Hz, 2H), 6.91(d, $J = 8.4$ Hz, 2H), 6.17(m, 1H), 4.50(m, 1H), 4.22(m, 1H), 3.72(m, 1H), 3.48(m, 2H), 3.30(m, 2H), 2.97(m, 1H), 2.53(m, 1H), 2.31(m, 1H), 2.00(m, 3H), 1.72(m, 3H), 1.50(m, 8H), 1.10(m, 4H), 0.83(m, 2H); LCMS: 527.5 ($M + H$)⁺.

[0115] Example 15. (*S*)-2-{2-[4-(4-Acetyl-piperazin-1-yl)-phenoxy]-acetyl-amino}-3-cyclohexyl-N-[2-(4-trifluoromethoxy-phenylamino)-ethyl]-propionamide; * $C_{32}H_{42}F_3N_5O_5$; ¹H NMR ($CDCl_3$) δ (ppm) 7.84(m, 1H), 7.26(m, 2H), 7.15(m, 2H), 7.11(m, 2H), 6.84(m, 2H),

4.38(m, 1H), 4.19(m, 1H), 3.85(m, 2H), 3.72(m, 2H), 3.51(m, 2H), 3.34(m, 1H), 3.20(m, 4H), 2.06(s, 3H), 1.54(m, 7H), 1.04(m, 6H), 0.80(m, 2H); LCMS: 634.5(M+H)⁺.

[0116] **Example 16.** (S)-2-Chloro-N-{1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-3-methyl-benzamide; * C₂₆H₂₈ClN₃O₃; LCMS: 466.2(M+H)⁺.

5 [0117] **Example 17.** 3-Cyclohexyl-2-[2-(4-methoxy-phenyl)-acetylamino]-N-[2-(4-trifluoromethoxy-phenylamino)-ethyl]-propionamide; * C₂₇H₃₄F₃N₃O₄; ¹H NMR (CDCl₃) δ(ppm)7.12((m, 1H), 7.04(m, 4H), 6.77(m, 4H), 5.81(m, 1H), 4.18(m, 1H), 3.71(s, 3H), 3.43(m, 4H), 3.23(m, 2H), 1.58(m, 7H), 1.35(m, 1H), 1.04(m, 3H), 1.35(m, 2H); LCMS: 522.5 (M+H)⁺.

10 [0118] **Example 18.** (S)-N-{2-[4-(3,5-Dichloro-phenoxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide.^{ss} Following the procedure of Example 327, except substituting phenyl-boronic acid for 3,5-dichlorophenyl-boronic acid in Step D, the title compound was prepared as a white solid (2 mg, 7%): ¹H-NMR (400MHz, CD₃OD) δ = 7.61-6.68 (m, 15H), 4.68 (dd, J= 7.1, J= 8.5, 1H), 3.79 (s, 3H), 3.57-3.10 (m, 15
6H), 2.38 (s, 3H). MS calcd. for C₃₂H₃₂Cl₂N₃O₄ (M+H⁺) 592.18, found 592.4.

[0119] **Example 19.** N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-methanesulfonyl-benzamide; * C₂₆H₃₂F₃N₃O₅S; ¹H NMR (CDCl₃) δ(ppm)7.87(m, 7H), 7.23(m, 3H), 4.41(m, 1H), 3.52(m, 3H), 3.34(m, 1H), 2.97(s, 3H), 1.68(m, 7H), 1.35(m, 1H), 1.11(m, 3H), 0.91(m, 2H); LCMS:556.4 (M+H)⁺.

20 [0120] **Example 20.** (S)-4-Benzyloxy-N-{1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-3,5-dimethyl-benzamide. * C₃₄H₃₇N₃O₄; LCMS: 552.25(M+H)⁺.

[0121] **Example 21.** (S)-4-Methoxy-N-{1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-3,5-dimethyl-benzamide. * C₂₈H₃₃N₃O₄; LCMS: 476.2(M+H)⁺.

25 [0122] **Example 22.** 5-Methoxy-1H-indole-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₈H₃₃F₃N₄O₄; LCMS: 547.5(M+H)⁺.

[0123] **Example 23.** 5-(3-Fluoro-phenyl)-furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide. * C₂₈H₃₁F₂N₃O₃; 496.5 (M+H)⁺.

[0124] **Example 24.** (S)-3-Bromo-N-{1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-4-methyl-benzamide; * C₂₆H₂₈BrN₃O₃; LCMS: 510.1(⁷⁹BrM+H)⁺, 512.1(⁸⁰BrM+H)⁺.

[0125] **Example 25.** Furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; *

C₂₃H₂₈F₃N₃O₄; ¹H NMR (CDCl₃) δ(ppm) 7.86(m, 1H), 7.65(m, 1H), 7.34(m, 2H), 7.25(m, 2H), 7.20(m, 1H), 7.04(m, 1H), 6.66(m, 1H), 4.60(m, 1H), 3.75(m, 2H), 3.60(m, 2H), 1.90(m, 7H), 1.55(m, 1H), 1.35(m, 3H), 1.11(m, 2H); LCMS: 468.4(M+H)⁺.

[0126] **Example 26.** Thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₃H₂₈F₃N₃O₃S; ¹H NMR (CDCl₃) δ(ppm) 7.64(m, 1H), 7.44(m, 2H), 7.07(m, 2H), 6.99(m, 3H), 6.78(m, 1H), 4.40(m, 1H), 3.50(m, 2H), 3.32(m, 2H), 1.62(m, 7H), 1.29(m, 1H), 1.09(m, 3H), 0.85(m, 2H); LCMS: 484.4(M+H)⁺.

[0127] **Example 27.** Furan-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₃H₂₈F₃N₃O₄; ¹H NMR (CDCl₃) δ(ppm) 7.84(m, 1H), 7.35(m, 2H), 7.04(d, J=8.0Hz, 2H), 6.83(m, 2H), 6.54(m, 2H), 4.40(m, 1H), 3.47(m, 2H), 3.28(m, 2H), 1.61(m, 7H), 1.26(m, 1H), 1.08(m, 3H), 0.85(m, 2H); LCMS: 468.4(M+H)⁺.

[0128] **Example 28.** N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-(5-methyl-[1,2,4]oxadiazol-3-yl)-benzamide. * C₂₈H₃₂F₃N₅O₄; LCMS: 560.5(M+H)⁺.

[0129] **Example 29.** 5-(4-Fluoro-phenyl)-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide. * C₂₉H₃₁F₄N₃O₃S; ¹H NMR (CDCl₃) δ(ppm) 7.55(m, 1H), 7.43(m, 3H), 7.05(m, 7H), 6.76(d, J=6.4Hz, 1H), 4.41(m, 1H), 3.51(m, 2H), 3.33(m, 2H), 1.61(m, 7H), 1.31(m, 1H), 1.13(m, 3H), 0.89(m, 2H); LCMS: 577.4 (M+H)⁺.

[0130] **Example 30.** (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-2,4,5-trimethyl-benzamide; * C₂₈H₃₃N₃O₃; LCMS: 460.2(M+H)⁺.

[0131] **Example 31.** 5-(3-Fluoro-phenyl)-furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide. * C₂₉H₃₁F₄N₃O₄; LCMS: 562.4 (M+H)⁺.

[0132] **Example 32.** 4-Benzyl-morpholine-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₀H₃₁F₃N₄O₄; ¹H NMR (CDCl₃) δ(ppm) 7.42(m, 4H), 7.35(m, 2H), 7.19(m, 2H), 7.04(m, 3H), 4.57(m, 1H), 4.33(m, 1H), 4.21(m, 1H), 4.11(m, 3H), 3.86(m, 1H), 3.59(m, 2H), 3.44(m, 2H), 3.34(m, 1H), 2.79(m, 2H), 1.73(m, 7H), 1.17(m, 4H), 0.93(m, 2H); LCMS: 577.5 (M+H)⁺.

[0133] **Example 33.** (S)-N-{2-[4-(4-Dimethylamino-phenoxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide.^{ss} Following the procedure of Example 327, except substituting phenyl-boronic acid for 4-dimethylaminophenyl-boronic acid in Step D, the title compound was prepared as a white solid (2 mg, 7%): MS calcd. for C₃₄H₃₉N₄O₄ (M+H)⁺ 567.30, found 567.5.

[0134] **Example 34.** 2'-Chloro-biphenyl-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide. * C₃₁H₃₃ClF₃N₃O₃; ¹H NMR (CD₃OD) δ(ppm) 7.93(m, 2H), 7.56(m, 3H), 7.41(m, 3H), 7.02(m, 2H), 6.76(m, 2H), 5.40(m, 1H), 3.44(m, 2H), 3.29(m, 2H), 1.74(m, 7H), 1.42(m, 1H), 1.19(m, 3H), 0.97(m, 2H); LCMS: 588.4(M+H)⁺.

[0135] **Example 35.** 5-(2-Trifluoromethyl-phenyl)-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₀H₃₁F₆N₃O₃S; ¹H NMR (CDCl₃) δ(ppm) 7.83(m, 1H), 7.69(d, J=7.6Hz, 1H), 7.41(m, 4H), 7.10(s, 4H), 6.99(m, 1H), 6.82(m, 1H), 4.38(m, 1H), 3.55(m, 2H), 3.39(m, 2H), 1.65(m, 7H), 1.34(m, 1H), 1.12(m, 3H), 0.89(m, 2H); LCMS: 628.4 (M+H)⁺.

[0136] **Example 36.** 5-(3-Fluoro-phenyl)-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₉H₃₁F₄N₃O₃S; ¹H NMR (CDCl₃) δ(ppm) 7.41(m, 1H), 7.28(m, 3H), 7.19(m, 2H), 7.04(d, J=8.0Hz, 2H), 7.11(m, 1H), 6.85(d, J=12.0Hz, 2H), 6.64(m, 1H), 4.43(m, 1H), 3.49(m, 2H), 3.30(m, 2H), 1.16(m, 7H), 1.32(m, 1H), 1.13(m, 3H), 0.88(m, 2H); LCMS: 578.3 (M+H)⁺.

[0137] **Example 37.** Thiophene-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₃H₂₈F₃N₃O₃S; ¹H NMR (CDCl₃) δ(ppm) 7.82(m, 1H), 7.41(m, 1H), 7.26(m, 2H), 7.01(m, 2H), 6.78(m, 3H), 4.46(m,

1H), 3.45(m, 2H), 3.27(m, 2H), 1.63(m, 7H), 1.29(m, 1H), 1.08(m, 3H), 0.87(m, 2H); LCMS: 484.4 (M+H)⁺.

[0138] Example 38. 5-Oxo-1-thiophen-2-ylmethyl-pyrrolidine-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; *

5 C₂₈H₃₅F₃N₄O₄S; ¹H NMR (CDCl₃) δ(ppm) 7.61(m, 1H), 7.11(m, 5H), 6.83(m, 2H), 6.51(m, 1H), 4.47(m, 2H), 4.18(m, 1H), 3.52(m, 6H), 3.06(m, 1H), 2.60(m, 2H), 1.60(m, 6H), 1.46(m, 1H), 1.09(m, 4H), 0.83(m, 2H); LCMS: 581.4 (M+H)⁺.

[0139] Example 39. 1-Furan-2-ylmethyl-5-oxo-pyrrolidine-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; *

10 C₂₈H₃₅F₃N₄O₅; ¹H NMR (CDCl₃) δ(ppm) 7.76(m, 1H), 7.18(m, 5H), 6.62(m, 1H), 6.16(m, 2H), 4.29(m, 3H), 3.41(m, 6H), 3.07(m, 1H), 2.61(m, 2H), 1.60(m, 7H), 1.08(m, 4H), 0.81(m, 2H); LCMS: 565.5 (M+H)⁺.

[0140] Example 40. 2-Methyl-5-(pyrrolidine-1-sulfonyl)-furan-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; *

15 C₂₈H₃₇F₃N₄O₆S; ¹H NMR (CDCl₃) δ(ppm) 7.37(m, 1H), 7.09(m, 3H), 6.97(m, 2H), 6.61(m, 1H), 4.33(m, 1H), 3.50(m, 3H), 3.27(m, 5H), 2.41(s, 3H), 1.77(m, 4H), 1.66(m, 7H), 1.29(m, 1H), 1.08(m, 3H), 0.86(m, 2H); LCMS: 615.4 (M+H)⁺.

[0141] Example 41. (S)-1-Phenyl-5-trifluoromethyl-1H-pyrazole-4-carboxylic acid {1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-amide; * C₂₉H₂₈F₃N₅O₃; ¹H

20 NMR (CD₃Cl) δ(ppm) 8.38(bs, 1H), 7.99(s, 1H), 7.71(m, 3H), 7.58(m, 4H), 7.52(m, 5H), 7.13(m, 3H), 4.84(dd, *J*=5.4Hz, *J*=13.9Hz, 1H), 4.03(s, 3H), 3.87(m, 1H), 3.66(m, 3H), 3.56(m, 1H), 3.37(m, 1H); LCMS: 552.2(M+H)⁺.

[0142] Example 42. 5-p-Tolyl-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₀H₃₄F₃N₃O₃S; ¹H NMR

25 (CDCl₃) δ(ppm) 7.40(m, 3H), 7.14(m, 3H), 6.94(d, *J*=8.4Hz, 2H), 6.79(m, 1H), 6.59(d, *J*=8.8Hz, 2H), 6.37(d, *J*=7.6Hz, 1H), 4.48(m, 1H), 3.45(m, 2H), 3.23(m, 2H), 2.30(s, 3H), 1.63(m, 7H), 1.29(m, 1H), 1.08(m, 3H), 0.88(m, 2H); LCMS: 574.5 (M+H)⁺.

[0143] Example 43. 4-Benzoimidazol-1-ylmethyl-N-{2-cyclohexyl-1-(S)-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide; * C₃₃H₃₆F₃N₅O₃S; ¹H

30 NMR (CDCl₃) δ(ppm) 9.41(s, 1H), 8.14(m, 1H), 7.82(d, *J*=8.0Hz, 1H), 7.45(m, 6H), 7.05(d, *J*=8.0Hz, 2H), 6.98(d, *J*=12Hz, 2H), 6.87(d, *J*=8.0Hz, 2H), 5.43(m, 2H), 4.53(m, 1H),

3.43(m, 2H), 3.26(m, 2H), 1.55(m, 7H), 1.29(m, 1H), 1.04(m, 3H), 0.84(m, 2H); LCMS: 608.5 (M+H)⁺.

[0144] Example 44. (S)-1-(4-Chloro-phenyl)-5-trifluoromethyl-1H-pyrazole-4-carboxylic acid {1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-amide; *

5 C₂₉H₂₇ClF₃N₅O₃; LCMS: 586.2(M+H)⁺.

[0145] Example 45. (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-p-tolyloxy-phenyl)-ethyl]-3-methyl-benzamide.^{ss} Following the procedure of Example 327, except substituting phenyl-boronic acid for 4-methylphenyl-boronic acid in Step D, the title compound was prepared as a white solid (6 mg, 22%): ¹H-NMR (400MHz, CD₃OD) δ =
10 7.60-6.79 (m, 16H), 4.63 (dd, J= 7.2, J= 8.3, 1H), 3.81 (s, 3H), 3.61-3.06 (m, 6H), 2.38 (s, 3H), 2.30 (s, 3H). MS calcd. for C₃₃H₃₆N₃O₄ (M+H⁺) 538.27, found 538.4.

[0146] Example 46. 3-Cyclohexyl-2-(S)-(2-tetrazol-1-yl-acetylamino)-N-[2-(4-trifluoromethoxy-phenylamino)-ethyl]-propionamide; * C₂₁H₂₈F₃N₇O₃; ¹H NMR (CDCl₃) δ(ppm) 8.81(s, 1H), 7.85(m, 1H), 7.56(m, 1H), 7.17(m, 4H), 5.16(m, 2H), 4.27(m, 1H),
15 3.50(m, 2H), 3.41(m, 1H), 3.33(m, 1H), 1.51(m, 7H), 1.09(m, 4H), 0.83(m, 2H); LCMS: 484.5 (M+H)⁺.

[0147] Example 47. 5-m-Tolyl-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₀H₃₄F₃N₃O₃S; ¹H NMR (CDCl₃) δ(ppm) 7.40(d, J=4.0Hz, 1H), 7.31(m, 2H), 7.20(m, 1H), 7.14(d, J=4Hz, 1H),
20 7.09(m, 1H), 7.02(m, 1H), 6.95(d, J=8Hz, 2H), 6.62(m, 2H), 6.54(d, J=7.2Hz, 1H), 4.49(m, 1H), 3.44(m, 2H), 3.22(m, 2H), 2.31(s, 3H), 1.63(m, 7H), 1.31(m, 1H), 1.11(m, 3H), 0.87(m, 2H); LCMS: 574.5 (M+H)⁺.

[0148] Example 48. 2,7-Dimethyl-pyrazolo[1,5-a]pyrimidine-6-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; *

25 C₂₇H₃₃F₃N₆O₃; ¹H NMR (CDCl₃) δ(ppm) 8.42(s, 1H), 7.38(m, 1H), 7.07(m, 2H), 6.93(m, 3H), 6.44(s, 1H), 4.48(m, 1H), 3.55(m, 2H), 3.37(m, 2H), 2.83(s, 3H), 2.45(s, 3H), 1.63(m, 7H), 1.31(m, 1H), 1.10(m, 3H), 0.90(m, 2H); LCMS: 547.5 (M+H)⁺.

[0149] Example 49. 2-Methyl-5-(morpholine-4-sulfonyl)-furan-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; *

30 C₂₈H₃₇F₃N₄O₇S; ¹H NMR (CDCl₃) δ(ppm) 7.61(m, 1H), 7.17(m, 3H), 7.07(m, 2H), 6.78(d,

$J=6.0\text{Hz}$, 1H), 4.33(m, 1H), 3.67(m, 4H), 3.48(m, 3H), 3.33(m, 1H), 3.10(m, 4H), 2.42(s, 3H), 1.63(m, 7H), 1.29(m, 1H), 1.19(m, 3H), 0.89(m, 2H); LMCS: 631.5(M+H)⁺.

[0150] Example 50. 5-(3-Trifluoromethyl-phenyl)-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; *

5 C₃₀H₃₁F₆N₃O₃S; ¹H NMR (CDCl₃) δ(ppm) 7.75(s, 1H), 7.67(d, $J=7.6\text{Hz}$, 1H), 7.53(d, $J=8.0\text{Hz}$, 1H), 7.45(m, 2H), 7.23(m, 1H), 7.05(m, 1H), 6.99(d, $J=8.0\text{Hz}$, 2H), 6.71(m, 2H), 6.65(d, $J=7.2\text{Hz}$, 1H), 4.48(m, 1H), 3.48(m, 2H), 3.27(m, 2H), 1.63(m, 7H), 1.32(m, 1H), 1.10(m, 3H), 0.90(m, 2H); LCMS: 628.4(M+H)⁺.

[0151] Example 51. 5-m-Tolyl-furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-

10 trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₀H₃₄F₃N₃O₄; ¹H NMR (CDCl₃) δ(ppm) 7.82(m, 1H), 7.43(m, 2H), 7.24(m, 1H), 7.12(m, 5H), 7.02(d, $J=3.6\text{Hz}$, 1H), 6.96(d, $J=6.0\text{Hz}$, 1H), 6.62(d, $J=3.6\text{Hz}$, 1H), 4.41(m, 1H), 3.56(m, 2H), 3.40(m, 2H), 2.33(s, 3H), 1.67(m, 7H), 1.34(m, 1H), 1.13(m, 3H), 0.90(m, 2H); LCMS: 558.5 (M+H)⁺.

[0152] Example 52. (S)-2,3-Dihydro-benzofuran-7-carboxylic acid {1-[2-(4-methoxy-

15 phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-amide; * C₂₇H₂₉N₃O₄; LCMS: 460.2(M+H)⁺.

[0153] Example 53. 5-Methanesulfonyl-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-

20 [2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₄H₃₀F₃N₃O₅S₂; ¹H NMR (CDCl₃) δ(ppm) 7.65(m, 1H), 7.55(m, 1H), 7.44(m, 1H), 7.23(m, 1H), 7.16(m, 2H), 7.06(m, 2H), 4.38(m, 1H), 3.51(m, 2H), 3.43(m, 1H), 3.33(m, 1H), 3.10(s, 3H), 1.61(m, 7H), 1.18(m, 1H), 0.89(m, 3H), 0.86(m, 2H); LCMS: 562.4 (M+H)⁺.

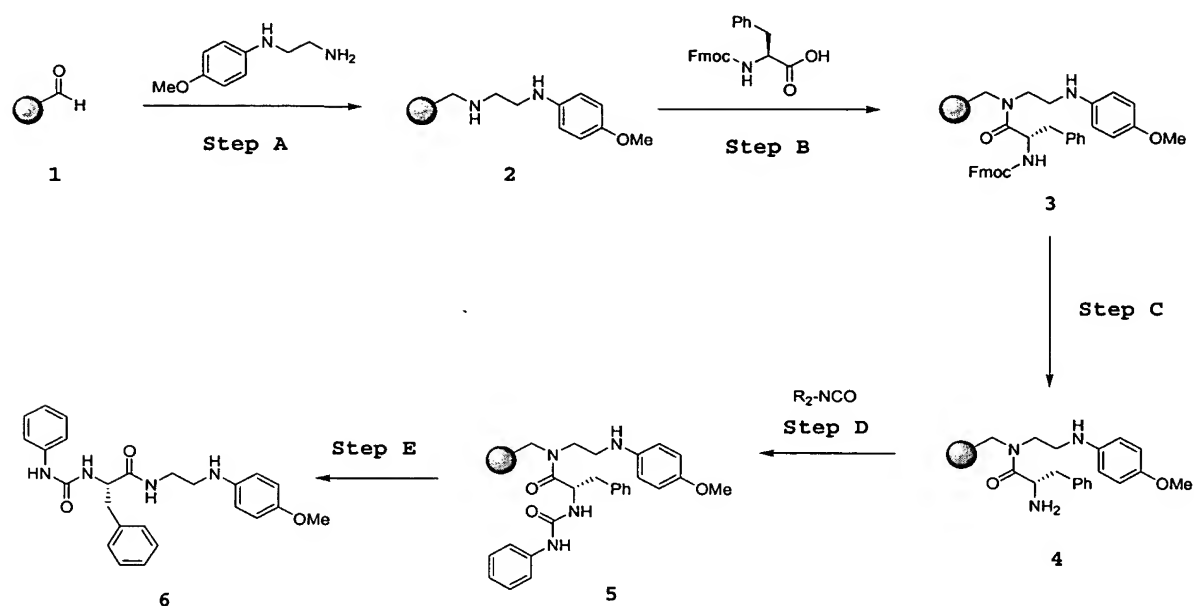
[0154] Example 54. 2-Phenyl-thiazole-4-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-

25 trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₈H₃₁F₃N₄O₃S; ¹H NMR (CDCl₃) δ(ppm) 7.91(s, 1H), 7.86(m, 2H), 7.78(d, $J=4\text{Hz}$, 1H), 7.51(m, 1H), 7.40(m, 3H), 7.06(m, 2H), 6.97(m, 2H), 4.44(m, 1H), 3.53(m, 2H), 3.35(m, 2H), 1.81(m, 1H), 1.67(m, 6H), 1.35(m, 1H), 1.14(m, 3H), 0.91(m, 2H); LCMS: 561.4 (M+H)⁺.

[0155] Example 55. (S)-3-Cyano-N-{1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-benzamide; * C₂₆H₂₆N₄O₃; LCMS: 443.2(M+H)⁺.

[0156] Example 56. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-3-(2-methyl-thiazol-4-yl)-benzamide; * C₂₉H₃₀N₄O₃S; LCMS: 515.2(M+H)⁺.

[0157] **Example 57.** (S)-N-[2-(4-Methoxy-phenylamino)-ethyl]-3-phenyl-2-(3-phenyl-ureido)-propionamide;



Solid phase synthesis: Steps A, B, C and E were described in Preparation 1.

- 5 **Step D:** Resin 4 (0.077 mmol) was swelled in DMF (1 mL) and phenyl isocyanate (18mg, 0.154 mmol, 2eq) was added in 1 mL DMF followed by pyridine (0.012 mL, 0.154 mmol, 2eq). The reaction was agitated for 3 h and then washed (DMF \times 4, then dichloromethane \times 3) and dried under nitrogen to yield resin 5. $C_{25}H_{28}N_4O_3$; 1H NMR (CD_3OD) δ (ppm) 7.27(m, 1H); 7.00(m, 3H); 4.38(m, 1H); 3.80(s, 3H); 3.61(m, 1H); 3.47(m, 1H); 3.38(m, 2H);
- 10 3.09(m, 1H); 2.99(m, 1H); LCMS: 433.2(M+H) $^+$. (14mg, 0.032mmol, 42%).

[0158] **Example 58.** 3-Cyclohexyl-2-(S)-(2-(S)-hydroxy-2-phenyl-acetyl-amino)-N-[2-(4-trifluoromethoxy-phenylamino)-ethyl]-propionamide; $C_{26}H_{32}F_3N_3O_4$; LCMS: 508.5 (M+H) $^+$.

- [0159] **Example 59.** Benzo[c]isoxazole-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; $C_{26}H_{29}F_3N_4O_4$; 1H NMR ($CDCl_3$) δ (ppm) 8.12(m, 1H), 7.50(m, 3H), 7.39(m, 2H), 7.17(m, 3H), 6.97(m, 1H), 4.46(m, 1H), 3.59(m, 4H), 1.65(m, 7H), 1.39(m, 1H), 1.16(m, 3H), 0.92(m, 2H); LCMS: 519.5 (M+H) $^+$.
- 15

- [0160] **Example 60.** N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-difluoromethoxy-benzamide; $C_{26}H_{30}F_3N_3O_4$; 1H NMR ($CDCl_3$) δ (ppm) 7.74(m, 3H), 7.11(m, 6H), 6.96(d, $J=6$ Hz, 1H), 6.55(t, $J=72.8$ Hz, 1H) 4.49(m, 1H),
- 20

3.59(m, 2H), 3.47(m, 1H), 3.39(m, 1H), 1.70(m, 7H), 1.37(m, 1H), 1.19(m, 3H), 0.97(m, 2H); LCMS: 544.4(M+H)⁺.

[0161] **Example 61.** N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-isopropoxy-benzamide; * C₂₈H₃₆F₃N₃O₄; ¹H NMR (CDCl₃) δ(ppm)
5 7.66(m, 3H), 7.14(d, J=8.4Hz, 2H), 7.02(d, J=8.8Hz, 2H), 6.87(d, J=9.2Hz, 2H), 6.70(d, J=6Hz, 1H), 4.61(m, 1H), 4.49(m, 1H), 3.58(m, 2H), 3.38(m, 2H), 1.70(m, 7H), 1.35(m, 7H), 1.19(m, 3H), 0.98(m, 2H); LCMS: 536.5(M+H)⁺.

[0162] **Example 62.** 5-Phenyl-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₉H₃₂F₃N₃O₃S; ¹H NMR
10 (CDCl₃) δ(ppm) 7.50(m, 2H), 7.42(m, 1H), 7.32(m, 4H), 7.17(m, 1H), 7.02(m, 2H), 6.84(m, 2H), 6.64(m, 1H), 4.43(m, 1H), 3.48(m, 2H), 3.29(m, 2H), 1.61(m, 7H), 1.29(m, 1H), 1.09(m, 3H), 0.89(m, 2H); LCMS: 560.3 (M+H)⁺.

[0163] **Example 63.** (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-nicotinamide. * C₂₄H₂₆N₄O₃; LCMS: 419.2(M+H)⁺.

15 [0164] **Example 64.** (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-isonicotinamide; * C₂₄H₂₆N₄O₃; LCMS: 419.2(M+H)⁺.

[0165] **Example 65.** 5-Phenyl-furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₉H₃₂F₃N₃O₄; ¹H NMR
20 (CDCl₃) δ(ppm) 7.75(m, 1H), 7.64(m, 2H), 7.35(m, 3H), 7.12(s, 3H), 7.02(d, J=3.6Hz, 1H), 6.93(m, 1H), 6.65(m, 2H), 4.39(m, 1H), 3.56(m, 2H), 3.39(m, 2H), 1.66(m, 7H), 1.35(m, 1H), 1.13(m, 3H), 0.92(m, 2H); LCMS: 544.4 (M+H)⁺.

[0166] **Example 66.** (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-o-tolyloxy-phenyl)-ethyl]-3-methyl-benzamide.^{ss} Following the procedure of Example 327, except substituting phenyl-boronic acid for 2-methylphenyl-boronic acid in Step D, the title
25 compound was prepared as a white solid (6 mg, 22%): ¹H-NMR (400MHz, CD₃OD) δ = 7.59-6.77 (m, 16H), 4.63 (dd, J= 7.1, J= 8.4, 1H), 3.81 (s, 3H), 3.61-3.05 (m, 6H), 2.37 (s, 3H), 2.14 (s, 3H). MS calcd. for C₃₃H₃₆N₃O₄ (M+H⁺) 538.27, found 538.4.

[0167] **Example 67.** N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-oxazol-5-yl-benzamide; *

C₂₈H₃₁F₃N₄O₄; ¹H NMR (CDCl₃) δ(ppm) 7.93(m, 1H), 7.73(d, J=8.0Hz, 2H), 7.63(d, J=8.0Hz, 2H), 7.39(s, 1H), 7.11(m, 1H), 7.01(d, J=8.0Hz, 2H), 6.76(m, 2H), 6.70(d, J=8.0Hz, 1H), 4.49(m, 1H), 3.49(m, 2H), 3.29(m, 2H), 1.65(m, 7H), 1.31(m, 1H), 1.10(m, 3H), 0.88(m, 2H); LCMS: 545.5(M+H)⁺.

5 [0168] **Example 68.** 5-(3-Trifluoromethyl-phenyl)-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;^{*}
C₃₀H₃₁F₆N₃O₃S; ¹H NMR (CDCl₃) δ(ppm) 7.92(m, 1H), 7.71(m, 1H), 7.64(d, J=8.0Hz, 1H),
7.52(m, 1H), 7.46(m, 2H), 7.22(m, 1H), 7.17(m, 4H), 6.99(d, J=6Hz, 1H), 4.39(m, 1H),
3.56(m, 2H), 3.40(m, 2H), 1.64(m, 7H), 1.34(m, 1H), 1.11(m, 3H), 0.89(m, 2H); LCMS:
10 628.4(M+H)⁺.

[0169] **Example 69.** 5-(2-Trifluoromethyl-phenyl)-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;^{*}
C₃₀H₃₁F₆N₃O₃S; ¹H NMR (CDCl₃) δ(ppm) 8.14(m, 1H), 7.69(m, 1H), 7.45(m, 3H), 7.36(m,
1H), 7.26(m, 2H), 7.17(m, 2H), 7.00(m, 2H), 4.36(m, 1H), 3.57(m, 2H), 3.44(m, 2H),
15 1.64(m, 7H), 1.35(m, 1H), 1.13(m, 3H), 0.89(m, 2H); LCMS: 628.3 (M+H)⁺.

[0170] **Example 70.** 5-p-Tolyl-furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;^{*} C₃₀H₃₄F₃N₃O₄; ¹H NMR
(CDCl₃) δ(ppm) 7.94(m, 1H), 7.40(m, 2H), 7.16(m, 2H), 7.06(m, 4H), 6.93(m, 1H), 6.89(m,
1H), 6.47(m, 1H), 4.27(m, 1H), 3.48(m, 2H), 3.34(m, 2H), 2.21(s, 3H), 1.61(m, 7H), 1.26(m,
20 1H), 1.03(m, 3H), 0.82(m, 2H); LCMS: 558.4 (M+H)⁺.

[0171] **Example 71.** N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-[(4,6-dimethyl-pyrimidin-2-yl)-methyl-amino]-benzamide;^{*}
C₃₂H₃₉F₃N₆O₃; LCMS: 613.5(M+H)⁺.

[0172] **Example 72.** 1-(4,6-Dimethyl-pyrimidin-2-yl)-piperidine-4-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide;^{*}
25 C₃₀H₄₁F₃N₆O₃; ¹H NMR (CDCl₃) δ(ppm) 7.33(m, 1H), 6.96(m, 1H), 6.91(d, J=8.8Hz, 2H),
6.82(d, J=8.8Hz, 2H), 6.18(s, 1H), 4.45(m, 2H), 4.09(m, 1H), 3.31(m, 2H), 3.10(m, 2H),
2.94(m, 2H), 2.36(m, 1H), 2.22(s, 6H), 1.72(m, 2H), 1.56(m, 2H), 1.41(m, 7H), 1.07(m, 1H),
1.01(m, 3H), 0.64(m, 2H); LCMS: 591.5 (M+H)⁺.

30 [0173] **Example 73.** N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-(4,6-dimethoxy-pyrimidin-2-yloxy)-benzamide;^{*} C₃₁H₃₆F₃N₅O₆; ¹H

NMR (CDCl₃) δ(ppm) 7.69(m, 1H), 7.50(m, 2H), 7.33(m, 2H), 7.07(m, 4H), 6.86(d, *J*=6Hz, 1H), 5.72(s, 1H), 4.39(m, 1H), 3.73(m, 6H), 3.52(m, 2H), 3.37(m, 2H), 1.62(m, 7H), 1.33(m, 1H), 1.11(m, 3H), 0.89(m, 2H); LCMS: 632.5(M+H)⁺.

[0174] **Example 74.** 3'-Methoxy-biphenyl-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₂H₃₆F₃N₃O₄; ¹H NMR (CDCl₃) δ(ppm) 7.98(m, 1H), 7.86(m, 1H), 7.66(m, 1H), 7.60(m, 1H), 7.41(m, 1H), 7.29(m, 1H), 7.13(m, 4H), 7.06(m, 1H), 7.03(m, 2H), 6.80(m, 1H), 4.43(m, 1H), 3.78(s, 3H), 3.54(m, 2H), 3.39(m, 2H), 1.68(m, 7H), 1.60(m, 1H), 1.10(m, 3H), 0.91(m, 2H); LCMS: 584.4 (M+H)⁺.

[0175] **Example 75.** N-{3-Cyclohexyl-1-(S)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-2-(R)-phenyl-butyramide; * C₂₉H₄₁N₃O₃; ¹H NMR (CDCl₃) δ(ppm) 7.89(m, 1H), 7.19(m, 3H), 7.13(m, 2H), 7.07(d, *J*=9.2Hz, 2H), 6.79(m, 2H), 6.25(m, 1H), 3.92(m, 1H), 3.73(s, 3H), 3.28(m, 5H), 1.96(m, 1H), 1.75(m, 2H), 1.53(m, 6H), 1.06(m, 6H), 0.75(m, 5H); LCMS: 480.6(M+H)⁺.

[0176] **Example 76.** 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-(2-thiophen-2-yl-acetyl-amino)-propionamide; * C₂₃H₃₀FN₃O₂S; ¹H NMR (CDCl₃) δ(ppm) 8.35(m, 1H), 7.35(m, 2H), 7.13(m, 1H), 7.07(m, 2H), 6.89(m, 2H), 6.39(d, *J*=4.0Hz, 1H), 4.03(m, 1H), 3.69(m, 5H), 3.33(m, 1H), 1.58(m, 7H), 1.10(m, 4H), 0.82(m, 2H); LCMS: 432.5(M+H)⁺.

[0177] **Example 77.** 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-(2-thiophen-3-yl-acetyl-amino)-propionamide; * C₂₃H₃₀FN₃O₂S; ¹H NMR (CDCl₃) δ(ppm) 8.36(m, 1H), 7.33(m, 2H), 7.22(m, 1H), 7.07(m, 3H), 6.92(m, 1H), 6.24(d, *J*=4.0Hz, 1H), 4.01(m, 1H), 3.54(m, 5H), 3.32(m, 1H), 1.58(m, 7H), 1.07(m, 4H), 0.80(m, 2H); LCMS: 432.5(M+H)⁺.

[0178] **Example 78.** (S)-3-Bromo-N-{1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-ethyl}-benzamide; * C₂₅H₂₆BrN₃O₃; LCMS: 496.1(⁷⁹BrM+H)⁺, 498.1(⁸⁰BrM+H)⁺.

[0179] **Example 79.** 1-Acetyl-piperidine-4-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide; C₂₅H₃₇FN₄O₃; ¹H NMR (CDCl₃) δ(ppm) 8.33(m, 1H), 7.42(m, 2H), 7.12(m, 2H), 6.40(m, 1H), 4.51(m, 3H), 4.09(m, 1H), 3.58(m, 3H), 3.35(m, 1H), 2.98(m, 1H), 2.42(m, 2H), 2.03(m, 3H), 1.51(m, 10H), 1.13(m, 4H), 0.88(m, 2H); LCMS: 461.6(M+H)⁺.

[0180] **Example 80.** N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-(4,6-dimethoxy-pyrimidin-2-yl)-benzamide; * C₃₁H₃₆F₃N₅O₅; ¹H NMR (CDCl₃) δ(ppm) 8.40(d, J=8.0Hz, 2H), 7.91(m, 1H), 7.71(d, J=8.0Hz, 2H), 7.13(m, 4H), 7.01(d, J=5.6Hz, 1H), 5.93(s, 1H), 3.96(s, 6H), 3.56(m, 2H), 3.40(m, 2H), 1.68(m, 7H),
5 1.35(m, 1H), 1.14(m, 3H), 0.89(m, 2H); LCMS: 616.5(M+H)⁺.

[0181] **Example 81.** 1-(5-Bromo-pyrimidin-2-yl)-piperidine-4-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₈H₃₆BrF₃N₆O₃; LCMS: 641.4(⁷⁹BrM+H)⁺, 643.4(⁸⁰BrM+H)⁺.

[0182] **Example 82.** (S)-2-(2-Cyclopent-2-enyl-acetylamino)-N-[2-(4-methoxy-phenylamino)-ethyl]-3-phenyl-propionamide; * C₂₅H₃₁N₃O₃; LCMS: 422.2(M+H)⁺
10

[0183] **Example 83.** 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(2-1H-indol-3-yl-acetylamino)-propionamide; * C₂₇H₃₃FN₄O₂; ¹H NMR (CDCl₃) δ(ppm) 8.19(m, 2H), 7.45(d, J=7.6Hz, 1H), 7.32(m, 3H), 7.16(m, 2H), 7.01(m, 3H), 6.30(d, J=4.0Hz, 1H), 3.99(m, 1H), 3.53(m, 5H), 3.27(m, 1H), 1.50(m, 5H), 1.30(m, 2H), 0.91(m, 3H), 0.65(m,
15 3H); LCMS: 465.5(M+H)⁺.

[0184] **Example 84.** N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-3-methanesulfonylamino-benzamide; * C₂₅H₂₅FN₄O₄S; ¹H NMR (CDCl₃) δ(ppm) 8.63(m, 1H), 8.36(m, 1H), 7.53(s, 1H), 7.40(m, 3H), 7.27(m, 1H), 7.22(m, 1H), 7.02(m, 2H), 4.48(m, 1H), 3.50(m, 4H), 2.86(s, 3H), 1.63(m, 7H), 1.32(m, 1H), 1.09(m, 3H), 0.89(m, 2H);
20 LCMS: 505.5(M+H)⁺.

[0185] **Example 85.** 5-(3-Trifluoromethyl-phenyl)-furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₉H₃₁F₄N₃O₃; ¹H NMR (CDCl₃) δ(ppm) 8.59(m, 1H), 7.88(m, 2H), 7.55(m, 4H), 7.34(m, 1H), 7.18(m, 2H), 7.08(d, J=3.6Hz, 1H), 6.79(d, J=3.6Hz, 1H), 4.42(m, 1H), 3.64(m, 4H), 1.77(m, 7H),
25 1.46(m, 1H), 1.22(m, 3H), 1.01(m, 2H); LCMS: 546.5(M+H)⁺.

[0186] **Example 86.** 3-Cyclohexyl-2-(S)-(2-(R,S)-fluoro-2-phenyl-acetylamino)-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide; * C₂₅H₃₁F₂N₃O₂; LCMS: 444.5(M+H)⁺.

[0187] **Example 87.** 3-Cyclohexyl-N-(S)-[2-(4-fluoro-phenylamino)-ethyl]-2-[2-(4-trifluoromethoxy-phenyl)-acetylamino]-propionamide; * C₂₆H₃₁F₄N₃O₃; ¹H NMR (CDCl₃)
30 δ(ppm) 8.44(m, 1H), 7.35(m, 2H), 7.24(m, 2H), 7.10(m, 4H), 6.43(d, J=4.0Hz, 1H), 4.11(m,

1H), 3.61(m, 5H), 3.37(m, 1H), 1.65(m, 7H), 1.20(m, 4H), 0.89(m, 2H); LCMS: 510.5(M+H)⁺.

[0188] **Example 88.** (S)-2-[3-(4-Chloro-phenyl)-ureido]-N-[2-(4-methoxy-phenylamino)-ethyl]-3-phenyl-propionamide; ^s C₂₅H₂₇ClN₄O₃; 467.2(³⁵ClM+H)⁺, 469.2(³⁷ClM+H)⁺.

5 [0189] **Example 89.** (S)-N-[2-(4-Methoxy-phenylamino)-ethyl]-2-[3-(4-phenoxy-phenyl)-ureido]-3-phenyl-propionamide; ^s C₃₁H₃₂N₄O₄; ¹H NMR (CD₃Cl) δ(ppm) 8.56(bs, 1H), 7.71(bs, 1H), 7.28(m, 7H), 7.19(m, 3H), 7.03(m, 3H), 6.88(m, 4H), 6.73(d, J=8.8Hz, 2H), 6.15(bs, 1H), 4.25(m, 1H), 3.77(s, 3H), 3.57(m, 2H), 3.33(m, 1H), 3.21(m, 2H), 2.95(m, 1H); LCMS: 525.2(M+H)⁺.

10 [0190] **Example 90.** (S)-N-[2-(4-Methoxy-phenylamino)-ethyl]-2-(3-phenethyl-ureido)-3-phenyl-propionamide; ^s C₂₇H₃₂N₄O₃; LCMS: 461.2(M+H)⁺.

[0191] **Example 91.** (S)-2-[3-(4-Fluoro-benzyl)-ureido]-N-[2-(4-methoxy-phenylamino)-ethyl]-3-phenyl-propionamide; ^s C₂₆H₂₉FN₄O₃; LCMS: 465.2(M+H)⁺.

15 [0192] **Example 92.** N-(S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-(4,6-dimethyl-pyrimidin-2-ylamino)-benzamide; ^{*} C₃₁H₃₇F₃N₆O₃; ¹H NMR (CDCl₃) δ(ppm) 12.30(s, 1H), 7.76(m, 2H), 7.64(m, 2H), 7.34(m, 1H), 7.01(m, 2H), 6.78(m, 3H), 6.58(s, 1H), 4.46(m, 1H), 3.48(m, 2H), 3.27(m, 2H), 2.50(s, 6H), 1.62(m, 7H), 1.31(m, 1H), 1.08(m, 3H), 0.86(m, 2H); LCMS: 599.5(M+H)⁺.

20 [0193] **Example 93.** 1-(5-Bromo-pyrimidin-2-yl)-piperidine-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; ^{*} C₂₈H₃₆BrF₃N₆O₃; ¹H NMR (CDCl₃) δ(ppm) 8.24(m, 2H), 7.89(m, 1H), 7.26(m, 2H), 7.18(m, 2H), 7.05(m, 1H), 4.15(m, 1H), 4.01(m, 1H), 3.79(m, 1H), 3.50(m, 3H), 3.34(m, 1H), 2.46(m, 1H), 1.84(m, 3H), 1.57(m, 9H), 1.18(m, 1H), 1.02(m, 3H), 0.87(m, 2H); LCMS: 641.4(⁷⁹BrM+H)⁺, 643.4(⁸⁰BrM+H)⁺.

25 [0194] **Example 94.** (S)-2-(3-Benzo[1,3] dioxol-5-yl-ureido)-N-[2-(4-methoxy-phenylamino)-ethyl]-3-phenyl-propionamide; ^s C₂₆H₂₈N₄O₅; LCMS: 477.2(M+H)⁺.

[0195] **Example 95.** 3-Cyclohexyl-2-(S)-[2-(2,5-difluorophenyl)-acetylamin]-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide; ^{*} C₂₅H₃₀F₃N₃O₂; ¹H NMR (CDCl₃) δ(ppm) 8.27(m, 1H), 7.26(m, 2H), 7.02(m, 2H), 6.85(m, 3H), 6.44(m, 1H), 4.08(m, 1H), 3.50(m, 5H), 3.32(m, 1H), 1.59(m, 7H), 1.13(m, 4H), 0.83(m, 2H); LCMS: 462.5(M+H)⁺.

30

[0196] **Example 96.** (S)-2-[3-(3-Fluoro-benzyl)-ureido]-N-[2-(4-methoxy-phenylamino)-ethyl]-3-phenyl-propionamide; $^{\text{S}} \text{C}_{26}\text{H}_{29}\text{FN}_4\text{O}_3$; LCMS: 465.2(M+H) $^{+}$.

[0197] **Example 97.** (S)-N-[2-(4-Methoxy-phenylamino)-ethyl]-3-phenyl-2-(3-o-tolyl-ureido)-propionamide; $^{\text{S}} \text{C}_{26}\text{H}_{30}\text{N}_4\text{O}_3$; LCMS: 447.2(M+H) $^{+}$.

5 [0198] **Example 98.** (S)-N-{2-[4-(3,4-Dichloro-phenoxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. $^{\text{SS}}$ Following the procedure of Example 327, except substituting phenyl-boronic acid for 3,4-dichlorophenyl-boronic acid in Step D, the title compound was prepared as a white solid (5 mg, 17%): ^1H -NMR (400MHz, CD_3OD) δ = 7.61-6.83 (m, 15H), 4.67 (dd, J = 7.0, J = 8.6, 1H), 3.81 (s, 3H), 3.61-3.09 (m, 10 6H), 2.38 (s, 3H). MS calcd. for $\text{C}_{32}\text{H}_{32}\text{Cl}_2\text{N}_3\text{O}_4$ (M+H) $^{+}$ 592.18, found 592.4.

[0199] **Example 99.** 3-Cyclohexyl-2-(S)-[2-(3,4-difluoro-phenyl)-acetylamino]-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide; $^{\text{*}} \text{C}_{25}\text{H}_{30}\text{F}_3\text{N}_3\text{O}_2$; ^1H NMR (CDCl_3) δ (ppm) 8.30(m, 1H), 7.25(m, 2H), 7.02(m, 5H), 6.35(m, 1H), 4.04(m, 1H), 3.51(m, 5H), 3.26(m, 1H), 1.63(m, 6H), 1.48(m, 1H), 1.13(m, 4H), 0.83(m, 2H); LCMS: 462.5 (M+H) $^{+}$.

15 [0200] **Example 100.** 3-Cyclohexyl-2-(S)-[2-(2,4-difluoro-phenyl)-acetylamino]-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide; $^{\text{*}} \text{C}_{25}\text{H}_{30}\text{F}_3\text{N}_3\text{O}_2$; ^1H NMR (CDCl_3) δ (ppm) 8.18(m, 1H), 7.12(m, 2H), 6.90(m, 3H), 6.50(m, 2H), 6.35(m, 1H), 3.90(m, 1H), 3.35(m, 5H), 3.17(m, 1H), 1.35(m, 7H), 0.96(m, 4H), 0.66(m, 2H); LCMS: 462.5(M+H) $^{+}$.

[0201] **Example 101.** (S)-N-[2-(4-Methoxy-phenylamino)-ethyl]-2-(3-naphthalen-1-yl-ureido)-3-phenyl-propionamide; $^{\text{S}} \text{C}_{29}\text{H}_{30}\text{N}_4\text{O}_3$; LCMS: 483.2(M+H) $^{+}$.

[0202] **Example 102.** (S)-2-[3-(2-tert-Butyl-6-methyl-phenyl)-ureido]-N-[2-(4-methoxy-phenylamino)-ethyl]-3-phenyl-propionamide; $^{\text{S}} \text{C}_{30}\text{H}_{38}\text{N}_4\text{O}_3$; LCMS: 503.3(M+H) $^{+}$.

[0203] **Example 103.** (S)-2-[3-(4-Acetyl-phenyl)-ureido]-N-[2-(4-methoxy-phenylamino)-ethyl]-3-phenyl-propionamide; $^{\text{S}} \text{C}_{27}\text{H}_{30}\text{N}_4\text{O}_4$; ^1H NMR (CD_3Cl) δ (ppm) 8.75(bs, 1H), 25 8.56(bs, 1H), 7.67(d, J =8.3Hz, 2H), 7.25(m, 10H), 6.88(d, J =8.6, 2H), 6.82(m, 1H), 4.37(m, 1H), 3.82(s, 3H), 3.64(m, 1H), 3.33(m, 3H), 3.15(m, 1H), 3.00(m, 1H), 2.47(s, 3H); LCMS: 475.2(M+H) $^{+}$.

[0204] **Example 104.** (S)-N-[2-(4-Methoxy-phenylamino)-ethyl]-2-[3-(3-methoxy-phenyl)-ureido]-3-phenyl-propionamide; $^{\text{S}} \text{C}_{26}\text{H}_{30}\text{N}_4\text{O}_4$; LCMS: 463.2(M+H) $^{+}$.

- [0205] Example 105.** (S)-Biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₁H₃₇N₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.72(m, 1H); 7.77(d, J=8.3Hz, 2H); 7.60(m, 2H), 7.58(m, 2H); 7.47(m, 5H); 7.16(m, 1H); 6.96(d, J=9.0Hz, 2H); 4.41(m, 1H); 3.82(s, 3H); 3.67(m, 3H); 3.50(m, 1H) 1.78(m, 7H); 1.49(m, 1H); 1.22(m, 3H); 1.00(m, 2H); LCMS: 500.25(M+H)⁺.
- [0206] Example 106.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-trifluoromethyl-benzamide; * C₂₆H₃₂F₃N₃O₃; LCMS: 492.4(M+H)⁺.
- [0207] Example 107.** 2-(S)-[2-(2-Chloro-4-fluoro-phenyl)-acetylamino]-3-cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide; * C₂₅H₃₀F₂N₃O₂; ¹H NMR (CDCl₃) δ(ppm) 8.31(m, 1H), 7.29(m, 2H), 7.22(m, 1H), 7.01(m, 3H), 6.85(m, 1H), 6.25(m, 1H), 4.06(m, 1H), 3.55(m, 5H), 3.32(m, 1H), 1.62(m, 6H), 1.46(m, 1H), 1.11(m, 4H), 0.83(m, 2H); LCMS: 478.5(M+H)⁺.
- [0208] Example 108.** (S)-2-Chloro-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide; * C₂₆H₃₄ClN₃O₃; LCMS: 472.2(M+H)⁺.
- [0209] Example 109.** (S)-4-Benzyloxy-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide; * C₃₂H₃₉N₃O₄; ¹H NMR (CD₃Cl) δ(ppm) 8.70(m, 1H), 7.64(d, J=8.9Hz, 2H), 7.45(d, J=8.9Hz, 2H), 7.38(m, 4H), 6.95(m, 5H), 6.84(m, 1H), 5.09(s, 2H), 4.33(m, 1H), 3.83(s, 3H), 3.64(m, 3H), 3.45(m, 1H), 1.75(m, 7H), 1.46(m, 1H), 1.25(m, 3H), 1.01(m, 2H); LCMS: 530.3(M+H)⁺.
- [0210] Example 110.** (S)-4-Benzyloxy-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3,5-dimethyl-benzamide; * C₃₄H₄₃N₃O₄; LCMS: 558.3(M+H)⁺.
- [0211] Example 111.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-methoxy-3,5-dimethyl-benzamide; * C₂₈H₃₉N₃O₄; LCMS: 482.3(M+H)⁺.
- [0212] Example 112.** (S)-3-Bromo-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-methyl-benzamide; * C₂₆H₃₄BrN₃O₃; LCMS: 516.4(⁷⁹BrM+H)⁺, 518.4(⁸⁰BrM+H)⁺.
- [0213] Example 113.** (S)-5-Fluoro-1H-indole-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide. * C₂₇H₃₃FN₄O₃; LCMS: 481.3(M+H)⁺.

[0214] **Example 114.** (S)-2-Amino-4-methyl-thiazole-5-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₃H₃₃N₅O₃S; LCMS: 460.2(M+H)⁺.

[0215] **Example 115.** (S)-1-Phenyl-5-trifluoromethyl-1H-pyrazole-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₉H₃₄F₃N₅O₃; LCMS: 558.2(M+H)⁺.

[0216] **Example 116.** (S)-1-(4-Chloro-phenyl)-5-trifluoromethyl-1H-pyrazole-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₉H₃₃ClF₃N₅O₃; LCMS: 592.2(M+H)⁺.

[0217] **Example 117.** (S)-5-(3-Trifluoromethyl-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₀H₃₄F₃N₃O₄; ¹H NMR (CD₃Cl) δ(ppm) 8.68(bs, 1H), 7.86(m, 2H), 7.56(m, 2H), 7.45(d, J=8.8Hz, 2H), 7.29(m, 1H), 7.06(d, J=3.3Hz, 1H), 6.95(d, J=8.7Hz, 2H), 6.77(d, J=3.5Hz, 1H), 4.40(m, 1H), 3.82(s, 3H), 3.69(m, 2H), 3.59(m, 1H), 3.52(m, 1H), 1.77(m, 7H), 1.47(m, 1H), 1.24(m, 3H), 1.00(m, 2H); LCMS: 558.2(M+H)⁺.

[0218] **Example 118.** (S)-3-Chloro-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide; * C₂₅H₃₂ClN₃O₃; LCMS: 458.2(³⁵ClM+H)⁺, 460.2(³⁷ClM+H)⁺.

[0219] **Example 119.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-dimethylamino-benzamide; * C₂₇H₃₈N₄O₃; LCMS: 467.3(M+H)⁺.

[0220] **Example 120.** (S)-3-Cyano-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide; * C₂₆H₃₂N₄O₃; LCMS: 449.2(M+H)⁺.

[0221] **Example 121.** (S)-4-Cyano-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide; * C₂₆H₃₂N₄O₃; LCMS: 449.2(M+H)⁺.

[0222] **Example 122.** N-{2-cyclohexyl-1-(S)-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-(R)-phenyl-propionamide; * C₂₇H₃₄F₃N₃O₃S **Error! Objects cannot be created from editing field codes.**; ¹H NMR (CDCl₃) δ(ppm) 7.18(m, 5H), 6.98(m, 2H), 6.79(m, 1H), 6.61(m, 2H), 5.73(d, J=7.2Hz, 1H), 4.20(m, 1H), 3.52(m, 1H), 3.31(m, 5H), 3.10(m, 2H), 1.58(m, 4H), 1.41(d, J=7.2Hz, 3H), 1.34(m, 1H), 1.04(m, 3H), 0.78(m, 2H); LCMS: 506.5 (M+H)⁺.

[0223] **Example 123.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-(2-methyl-thiazol-4-yl)-benzamide; * C₂₉H₃₆N₄O₃S; LCMS: 521.2(M+H)⁺.

[0224] **Example 124.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-[1,2,4]triazol-1-yl-benzamide; * C₂₇H₃₄N₆O₃; LCMS: 491.3(M+H)⁺.

[0225] **Example 125.** 3-Cyclohexyl-2-(S)-[2-(3,5-difluoro-phenyl)-acetylamino]-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide; * C₂₅H₃₀F₃N₃O₂; ¹H NMR (CDCl₃) δ(ppm) 8.40(m, 1H), 7.36(m, 2H), 7.12(m, 2H), 6.77(m, 2H), 6.66(m, 1H), 6.50(m, 1H), 4.13(m, 1H), 3.58(m, 5H), 3.36(m, 1H), 1.65(m, 7H), 1.18(m, 4H), 0.92(m, 2H); LCMS: 462.5(M+H)⁺.

[0226] **Example 126.** (S)-N-{3-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-3-trifluoromethyl-benzamide; * C₂₇H₃₄F₃N₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.75(bs, 1H), 8.06(bs, 1H), 7.97(d, J=7.6Hz, 1H), 7.78(d, J=7.8Hz, 1H), 7.58(m, 1H), 7.44(m, 3H), 6.95(d, J=8.5Hz, 2H), 4.34(m, 1H), 3.84(s, 3H), 3.71(m, 1H), 3.58(m, 3H), 1.99(m, 1H), 1.87(m, 1H), 1.69(m, 5H), 1.32(m, 6H), 0.89(m, 2H); LCMS: 506.4(M+H)⁺.

[0227] **Example 127.** (S)-N-{3-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-4-morpholin-4-yl-benzamide; * C₃₀H₄₂N₄O₄; ¹H NMR (CD₃Cl) δ(ppm) 8.70(bs, 1H), 7.65(d, J=8.6Hz, 2H), 7.49(d, J=8.6Hz, 2H), 7.07(m, 1H), 6.97(d, J=8.6Hz, 2H), 6.84(d, J=8.6Hz, 2H), 4.24(m, 1H), 3.85(m, 7H), 3.58(m, 2H), 3.54(m, 2H), 3.25(m, 4H), 1.96(m, 1H), 1.83(m, 1H), 1.69(m, 5H), 1.27(m, 6H), 0.89(m, 2H); LCMS: 523.5(M+H)⁺.

[0228] **Example 128.** (4-Cyclohexyl-N-[2-(4-methoxy-phenylamino)-ethyl]-2-(S)-(2-(S)-phenyl-propionylamino)-butyramide; * C₂₈H₃₉N₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.46(bs, 1H), 7.31(m, 7H), 6.90(m, 2H), 6.43(m, 1H), 3.95(m, 1H), 3.84(s, 3H), 3.68(m, 2H), 3.50(m, 2H), 3.34(m, 1H), 1.82(m, 1H), 1.66(m, 6H), 1.49(d, J=7.2Hz, 3H), 1.14(m, 6H), 0.82(m, 2H); LCMS: 466.5(M+H)⁺.

[0229] **Example 129.** (S)-4-Benzyloxy-N-{3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-benzamide; * C₃₃H₄₁N₃O₄; ¹H NMR (CD₃Cl) δ(ppm) 8.71(bs, 1H), 7.71(d, J=8.5Hz, 2H), 7.39(m, 7H), 7.17(m, 1H), 6.95(m, 4H), 5.10(s, 2H), 4.27(m, 1H),m

3.83(s, 3H), 3.66(m, 2H), 3.53(m, 2H), 1.93(m, 1H), 1.82(m, 1H), 1.68(m, 5H), 1.25(m, 6H), 0.87(m, 2H); LCMS: 544.5(M+H)⁺.

[0230] Example 130. (S)-Biphenyl-4-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide; * C₃₂H₃₉N₃O₃; ¹H NMR (CD₃Cl) δ(ppm)

5 8.80(bs, 1H), 7.81(d, *J*=7.9Hz, 2H), 7.60(m, 4H), 7.47(m, 5H), 7.19(m, 1H), 6.96(d, *J*=8.7Hz, 2H), 4.32(m, 1H), 3.84(s, 3H), 3.70(m, 2H), 3.57(m, 2H), 1.56(m, 1H), 1.43(m, 1H), 1.70(m, 5H), 1.22(m, 6H), 0.90(m, 2H); LCMS: 514.5(M+H)⁺.

[0231] Example 131. (S)-5-Chloro-1H-indole-2-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide; * C₂₈H₃₅ClN₄O₃; ¹H NMR (CD₃Cl)

10 δ(ppm) 10.19(bs, 1H), 8.69(bs, 1H), 7.37(m, 4H), 7.18(m, 2H), 6.90(m, 2H), 6.65(bs, 1H), 4.40(m, 1H), 3.80(s, 3H), 3.56(m, 4H), 1.84(m, 1H), 1.63(m, 6H), 1.19(m, 6H), 0.81(m, 2H); LCMS: 511.5(M+H)⁺.

[0232] Example 132. (S)-5-Fluoro-1H-indole-2-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide; * C₂₈H₃₅FN₄O₃; ¹H NMR (CD₃Cl)

15 δ(ppm) 10.03(bs, 1H), 8.66(bs, 1H), 7.42(d, *J*=8.5Hz, 2H), 7.26(m, 1H), 7.04(m, 2H), 6.90(m, 2H), 6.72(m, 1H), 4.39(m, 1H), 3.80(s, 3H), 3.52(m, 4H), 1.87(m, 1H), 1.72(m, 6H), 1.17(m, 6H), 0.81(m, 2H); LCMS: 495.5(M+H)⁺.

[0233] Example 133. (S)-2-Amino-4-methyl-thiazole-5-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide; * C₂₄H₃₅N₅O₃S; ¹H NMR

20 (CD₃Cl) δ(ppm) 9.61(bs, 2H), 8.30(bs, 1H), 7.55(m, 1H), 7.39(d, *J*=8.6Hz, 2H), 6.95(d, *J*=8.6Hz, 2H), 4.27(m, 1H), 3.83(s, 3H), 3.71(m, 1H), 3.51(m, 3H), 2.44(s, 3H), 1.93(m, 1H), 1.82(m, 1H), 1.68(m, 5H), 1.22(m, 6H), 0.87(m, 2H); LCMS: 474.5(M+H)⁺.

[0234] Example 134. (S)-5-Chloro-benzofuran-2-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide; * C₂₈H₃₄ClN₃O₄; ¹H NMR (CD₃Cl)

25 δ(ppm) 8.69(bs, 1H), 7.55(s, 1H), 7.40(m, 6H), 6.95(d, *J*=8.6Hz, 2H), 4.39(m, 1H), 3.83(s, 3H), 3.62(m, 4H), 1.99(m, 1H), 1.85(m, 1H), 1.69(m, 5H), 1.28(m, 6H), 0.89(m, 2H); LCMS: 512.4(M+H)⁺.

[0235] Example 135. N-{2-cyclohexyl-1-(S)-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-(R)-phenyl-butylamide; * C₂₈H₃₆F₃N₃O₃; ¹H NMR (CDCl₃) δ(ppm)

30 7.65(m, 1H), 7.08(m, 7H), 7.00(m, 2H), 6.37(d, *J*=5.6Hz, 1H), 4.14(m, 1H), 3.40(m, 2H),

3.21(m, 3H), 1.95(m, 1H), 1.74(m, 1H), 1.55(m, 7H), 1.44(m, 1H), 1.03(m, 3H), 0.79(m, 5H); LCMS: 520.5(M+H)⁺.

[0236] Example 136. (S)-5-(3-Trifluoromethyl-phenyl)-furan-2-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide; * C₃₁H₃₆F₃N₃O₄;

¹H NMR (CD₃Cl) δ(ppm) 8.77(bs, 1H), 7.89(m, 2H), 7.59(m, 2H), 7.48(m, 2H), 7.25(m, 1H), 7.11(d, *J*=3.5Hz, 1H), 6.97(d, *J*=8.7Hz, 2H), 6.97(d, *J*=3.4Hz, 1H), 4.29(m, 1H), 3.84(s, 3H), 3.73(m, 2H), 3.59(m, 2H), 2.01(m, 1H), 1.88(m, 1H), 1.74(m, 5H), 1.23(m, 6H), 0.91(m, 2H); LCMS: 572.5(M+H)⁺.

[0237] Example 137. (S)-Benzothiazole-6-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide; * C₂₇H₃₄N₄O₃S; ¹H NMR (CD₃Cl) δ(ppm) 9.15(s, 1H), 8.74(bs, 1H), 8.43(s, 1H), 8.12(d, *J*=8.5, 1H), 7.90(d, *J*=8.7, 1H), 7.60(m, 1H), 7.45(d, *J*=8.2, 2H), 6.94(d, *J*=8.2, 2H), 4.40(m, 1H), 3.83(s, 3H), 3.71(m, 1H), 3.65(m, 1H), 3.57(m, 2H), 1.96(m, 1H), 1.85(m, 1H), 1.68(m, 5H), 1.18(m, 6H), 0.88(m, 2H); LCMS: 495.5(M+H)⁺.

[0238] Example 138. (S)-N-{3-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-3-trifluoromethoxy-benzamide; * C₂₇H₃₄F₃N₃O₄; ¹H NMR (CD₃Cl) δ(ppm) 8.73(bs, 1H), 7.72(d, *J*=7.5Hz, 1H), 7.65(s, 1H), 7.43(m, 5H), 6.94(d, *J*=8.2Hz, 2H), 4.34(m, 1H), 3.83(s, 3H), 3.62(m, 4H), 1.94(m, 1H), 1.83(m, 1H), 1.68(m, 5H), 1.27(m, 6H), 0.88(m, 2H); LCMS: 522.5(M+H)⁺.

[0239] Example 139. (S)-3-Cyano-N-{3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-benzamide; * C₂₇H₃₄N₄O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.71(m, 1H), 8.07(s, 1H), 8.00(d, *J*=7.7Hz, 1H), 7.80(d, *J*=7.8Hz, 1H), 7.56(m, 1H), 7.43(d, *J*=8.7Hz, 2H), 7.35(d, *J*=4.3Hz, 1H), 6.98(d, *J*=8.7Hz, 2H), 4.33(m, 1H), 3.85(s, 3H), 3.62(m, 4H), 2.00(m, 1H), 1.86(m, 1H), 1.70(m, 5H), 1.26(m, 6H), 0.92(m, 2H); LCMS: 463.5(M+H)⁺.

[0240] Example 140. (S)-4-Cyano-N-{3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-benzamide; * C₂₇H₃₄N₄O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.67(bs, 1H), 7.88(d, *J*=8.1Hz, 2H), 7.71(d, *J*=8.1Hz, 2H), 7.45(m, 1H), 7.42(d, *J*=8.6Hz, 2H), 6.96(d, *J*=8.6Hz, 2H), 4.34(m, 1H), 3.85(s, 3H), 3.58(m, 4H), 1.98(m, 1H), 1.84(m, 1H), 1.70(m, 5H), 1.27(m, 6H), 0.89(m, 2H); LCMS: 463.5(M+H)⁺.

[0241] Example 141. N-{2-cyclohexyl-1-(S)-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-phenoxy-benzamide; * C₃₁H₃₄F₃N₃O₄; LCMS: 570.5 (M+H)⁺.

[0242] **Example 142.** (S)-N-{3-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-3-(2-methyl-thiazol-4-yl)-benzamide; * C₃₀H₃₈N₄O₃S; ¹H NMR (CD₃Cl) δ(ppm) 8.69(bs, 1H), 8.30(bs, 1H), 7.95(d, J=7.7Hz, 1H), 7.75(m, 2H), 7.46(m, 3H) 6.94(d, J=8.6Hz, 2H), 7.38(m, 1H), 4.35(m, 1H), 3.82(s, 3H), 3.59(m, 4H), 2.82(s, 3H),
5 1.92(m, 2H), 1.68(m, 5H), 1.22(m, 6H), 0.88(m, 2H); LCMS: 535.5(M+H)⁺.

[0243] **Example 143.** (S)-N-{3-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-4-[1,2,4]triazol-1-yl-benzamide; * C₂₈H₃₆N₆O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.72(bs, 2H), 8.16(bs, 1H), 7.92(d, J=8.3Hz, 2H), 7.74(d, J=8.3Hz, 2H), 7.56(d, J=5.3Hz, 1H), 7.45(d, J=8.6Hz, 2H), 6.96(d, J=8.6Hz, 2H), 4.36(m, 1H), 3.84(s, 3H),
10 3.70(m, 2H), 3.55(m, 2H), 1.99(m, 1H), 1.87(m, 1H), 1.68(m, 5H), 1.27(m, 6H), 0.89(m, 2H); LCMS: 505.5(M+H)⁺.

[0244] **Example 144.** (S)-Biphenyl-3-carboxylic acid {3-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-amide; * C₃₂H₃₉N₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.79(bs, 1H), 7.98(s, 1H), 7.74(d, J=7.6Hz, 2H), 7.45(m, 9H), 6.91(d, J=8.7Hz, 2H), 4.33(m,
15 1H), 3.80(s, 3H), 3.62(m, 4H), 1.97(m, 1H), 1.86(m, 1H), 1.68(m, 5H), 1.27(m, 6H), 0.88(m, 2H); LCMS: 514.5(M+H)⁺.

[0245] **Example 145.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-fluoro-benzamide; * C₂₅H₃₂FN₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.58(bs, 1H), 7.71(m, 1H), 7.44(m, 3H), 7.33(m, 1H), 7.13(m, 2H), 6.92(d, J=8.6Hz, 2H),
20 4.31(m, 1H), 3.79(s, 3H), 3.61(m, 2H), 3.46(m, 2H), 1.72(m, 7H), 1.41(m, 1H), 1.19(m, 3H), 0.96(m, 2H); LCMS: 442.4(M+H)⁺.

[0246] **Example 146.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3,4-difluoro-benzamide; * C₂₅H₃₁F₂N₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.68(bs, 1H), 7.57(m, 2H), 7.44(m, 3H), 7.21(m, 1H), 6.97(d, J=8.6Hz, 2H), 4.44(m, 1H),
25 3.85(s, 3H), 3.64(m, 3H), 3.48(m, 1H), 1.77(m, 7H), 1.45(m, 1H), 1.20(m, 3H), 0.96(m, 2H); LCMS: 460.5(M+H)⁺.

[0247] **Example 147.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-fluoro-2-methyl-benzamide; * C₂₆H₃₄FN₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.51(bs, 1H), 7.32(d, J=8.3Hz, 2H), 7.09(m, 3H), 6.82(d, J=8.4Hz, 2H), 6.58(m,
30 1H), 4.36(m, 1H), 3.72(s, 3H), 3.68(m, 1H), 3.45(m, 3H), 2.18(s, 3H), 1.62(m, 7H), 1.31(m, 1H), 1.16(m, 3H), 0.89(m, 2H); LCMS: 456.5(M+H)⁺.

[0248] **Example 148.** (S)-2-Chloro-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-5-methyl-benzamide; * C₂₆H₃₄ClN₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.58(bs, 1H), 7.53(d, J=8.5Hz, 2H), 7.43(s, 1H), 7.33(m, 1H), 7.24(m, 2H), 6.99(d, J=8.5Hz, 2H), 4.49(m, 1H), 3.87(s, 3H), 3.67(m, 4H), 2.33(s, 3H), 1.85(m, 7H), 1.53(m, 1H), 1.27(m, 3H), 1.03(m, 2H); LCMS: 472.4(³⁵ClM+H)⁺, 474.4(³⁷ClM+H)⁺.

[0249] **Example 149.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-fluoro-3-trifluoromethyl-benzamide; * C₂₆H₃₁F₄N₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.68(bs, 1H), 8.09(m, 1H), 8.03(m, 1H), 7.80(m, 1H), 7.41(d, J=8.0, 2H), 7.23(m, 1H), 6.95(d, J=7.9, 2H), 4.49(m, 1H), 3.84(s, 3H), 3.63(m, 3H), 3.48(m, 1H), 1.74(m, 7H), 1.46(m, 1H), 1.22(m, 3H), 0.98(m, 2H); LCMS: 510.4(M+H)⁺.

[0250] **Example 150.** (S)-5-Methyl-1-phenyl-1H-pyrazole-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₉H₃₇N₅O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.71(bs, 1H), 7.89 (bs, 1H), 7.44(m, 5H), 7.28(m, 2H), 6.90(m, 3H), 4.30(m, 1H), 3.78(s, 3H), 3.65(m, 3H), 3.40(m, 1H), 2.21(s, 3H), 1.70(m, 7H), 1.45(m, 1H), 1.20(m, 3H), 0.98(m, 2H); LCMS: 504.5(M+H)⁺.

[0251] **Example 151.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-propyl-benzamide; * C₂₈H₃₉N₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.69(bs, 1H), 7.66(d, J=7.8Hz, 2H), 7.47(d, J=8.6Hz, 2H), 7.30(m, 1H), 7.21(d, J=7.8Hz, 2H), 6.95(d, J=8.5Hz, 2H), 4.43(m, 1H), 3.84(s, 3H), 3.63(m, 3H), 3.48(m, 1H), 2.62(m, 2H), 1.72(m, 10H), ?(m, 1H), 1.22(m, 3H), 0.94(m, 4H); LCMS: 466.5(M+H)⁺.

[0252] **Example 152.** 3-Cyclohexyl-2-(S)-[2-(4-fluoro-phenyl)-acetylamino]-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide; * C₂₅H₃₁F₂N₃O₂; ¹H NMR (CDCl₃) δ(ppm) 8.44(m, 1H), 7.34(m, 2H), 7.19(m, 2H), 7.11(m, 2H), 6.94(m, 2H), 6.42(m, 1H), 4.11(m, 1H), 3.62(m, 5H), 3.39(m, 1H), 1.66(m, 6H), 1.32(m, 1H), 1.18(m, 4H), 0.89(m, 2H); LCMS: 444.5(M+H)⁺.

[0253] **Example 153.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-methoxy-benzamide; * C₂₆H₃₅N₃O₄; ¹H NMR (CD₃Cl) δ(ppm) 8.66(bs, 1H), 7.72(d, J=8.4Hz, 2H), 7.46(d, J=8.6Hz, 2H), 7.34(m, 1H), 6.96(d, J=8.5Hz, 2H), 6.88(d, J=8.4Hz, 2H), 4.43(dd, J=6.1Hz, J=13.6Hz, 1H), 3.84(s, 6H), 3.60(m, 3H), 3.48(m, 1H), 1.71(m, 7H), 1.46(m, 1H), 1.23(m, 3H), 0.96(m, 2H); LCMS: 454.5(M+H)⁺.

[0254] **Example 154.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-fluoro-5-trifluoromethyl-benzamide; * C₂₆H₃₁F₄N₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.66(bs, 1H), 8.13(d, *J*=6.5Hz, 1H), 7.79(m, 1H), 7.46(m, 3H), 7.33(m, 1H), 6.98(d, *J*=8.4Hz, 2H), 4.44(m, 1H), 3.85(s, 3H), 3.73(m, 2H), 3.62(m, 2H), 1.75(m, 7H), 1.47(m, 1H), 1.25(m, 3H), 1.04(m, 2H); LCMS: 510.5(M+H)⁺.

[0255] **Example 155.** (S)-3-Chloro-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-fluoro-benzamide; * C₂₅H₃₁ClFN₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.60(bs, 1H), 7.68(m, 1H), 7.58(m, 1H), 7.47(d, *J*=8.6Hz, 2H), 7.37(m, 1H), 7.15(m, 1H), 6.98(d, *J*=8.6Hz, 2H), 4.43(m, 1H), 3.85(s, 3H), 3.66(m, 3H), 3.55(m, 1H), 1.78(m, 7H), 1.47(m, 1H), 1.24(m, 3H), 1.04(m, 2H); LCMS: 476.4(M+H)⁺.

[0256] **Example 156.** (S)-5-Chloro-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-fluoro-benzamide; * C₂₅H₃₁ClFN₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.55(bs, 1H), 7.69(m, 1H), 7.35(m, 3H), 7.24(m, 1H), 7.03(m, 1H), 6.89(d, *J*=8.5HZ, 2H), 4.32(m, 1H), 3.75(s, 3H), 3.63(m, 2H), 3.49(m, 2H), 1.66(m, 7H), 1.35(m, 1H), 1.18(m, 3H), 0.92(m, 2H); 476.4(³⁵CIM+H)⁺, 478.4(³⁷CIM+H)⁺.

[0257] **Example 157.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-5-fluoro-2-methyl-benzamide; * C₂₆H₃₄FN₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.75(bs, 1H), 7.61(d, *J*=8.5Hz, 2H), 7.36(m, 2H), 7.24(m, 1H), 7.11(d, *J*=8.4Hz, 2H), 6.88(m, 1H), 4.63(m, 1H), 4.00(s, 3H), 3.98(m, 1H), 3.75(m, 3H), 2.53(s, 3H), 1.93(m, 7H), 1.60(m, 1H), 1.42(m, 3H), 1.18(m, 2H); LCMS: 456.5(M+H)⁺.

[0258] **Example 158.** (S)-1-Phenyl-cyclopropanecarboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₈H₃₇N₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.49(bs, 1H), 7.42(m, 7H), 7.00(d, *J*=8.7Hz, 2H), 4.01(m, 1H), 3.86(s, 3H), 3.67(m, 3H), 3.48(m, 1H), 1.60(m, 9H), 1.30(m, 1H), 1.21(m, 5H), 0.82(m, 3H); LCMS: 464.5(M+H)⁺.

[0259] **Example 159.** (S)-3-Cyclohexyl-N-[2-(4-methoxy-phenylamino)-ethyl]-2-(2-phenylamino-acetyl-amino)-propionamide; * C₂₆H₃₆N₄O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.31(bs, 1H), 7.79(m, 1H), 7.35(d, *J*=8.2Hz, 2H), 7.24(m, 2H), 6.95(m, 1H), 6.88(d, *J*=8.3Hz, 2H), 6.80(d, *J*=7.9Hz, 2H), 4.20(m, 1H), 3.86(m, 2H), 3.75(s, 3H), 3.55(m, 3H), 3.37(m, 1H), 1.52(m, 7H), 0.95(m, 4H), 0.76(m, 2H); LCMS: 453.5(M+H)⁺.

[0260] Example 160. 3-Cyclohexyl-2-(S)-(2-(R)-hydroxy-2-phenyl-acetyl-amino)-N-[2-(4-methoxy-phenyl-amino)-ethyl]-propionamide; * C₂₆H₃₅N₃O₄; ¹H NMR (CD₃Cl) δ(ppm) 8.42(bs, 1H), 7.50(m, 1H), 7.41(m, 2H), 7.32(m, 3H), 7.19(d, J=8.6Hz, 2H), 6.85(d, J=8.5Hz, 2H), 5.00(s, 1H), 4.17(m, 1H), 3.83(s, 3H), 3.73(m, 1H), 3.56(m, 2H), 3.31(m, 1H), 1.63(m, 7H), 1.10(m, 4H); 0.89(m, 2H).

[0261] Example 161. (S)-1-(4-Fluoro-phenyl)-5-methyl-1H-pyrazole-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenyl-amino)-ethylcarbamoyl]-ethyl}-amide; * C₂₉H₃₆FN₅O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.76(bs, 1H), 7.94(bs, 1H), 7.47(d, J=8.7Hz, 2H), 7.31(m, 2H), 7.17(m, 2H), 6.96(m, 3H), 4.36(m, 1H), 3.83(s, 3H), 3.73(m, 3H), 3.44(m, 1H): 2.24(s, 3H), 1.75(m, 7H), 1.49(m, 1H), 1.24(m, 3H), 1.01(m, 2H); LCMS: 522.5(M+H)⁺.

[0262] Example 162. (S)-1-(4-Methoxy-phenyl)-5-methyl-1H-pyrazole-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenyl-amino)-ethylcarbamoyl]-ethyl}-amide; * C₃₀H₃₉N₅O₄; LCMS: 534.5(M+H)⁺.

[0263] Example 163. (S)-1-(4-Chloro-phenyl)-5-methyl-1H-pyrazole-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenyl-amino)-ethylcarbamoyl]-ethyl}-amide; * C₂₉H₃₆ClN₅O₃; ¹H NMR (CD₃Cl) δ(ppm) 7.81(m, 2H), 7.36(m, 2H), 7.22(m, 2H), 7.01(m, 2H), 6.77(m, 2H), 6.53(m, 1H), 4.35(m, 1H), 3.69(s, 3H), 3.51(m, 3H), 3.27(m, 1H), 2.29(s, 3H), 1.69(m, 7H), 1.34(m, 1H), 1.10(m, 3H), 0.90(m, 2H); LCMS: 538.4(M+H)⁺.

[0264] Example 164. N-{2-Cyclohexyl-1-(S)-[2-(4-methoxy-phenyl-amino)-ethylcarbamoyl]-ethyl}-2-(S)-phenyl-butyramide; * C₂₈H₃₉N₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.34(m, 1H), 7.33(m, 2H), 7.25(m, 5H), 6.94(m, 2H), 6.53(m, 1H), 4.13(m, 1H), 3.80(s, 3H), 3.57(m, 2H), 3.40(m, 3H), 2.05(m, 1H), 1.84(m, 1H), 1.67(m, 7H), 1.15(m, 4H), 0.86(m, 5H); LCMS: 466.5(M+H)⁺.

[0265] Example 165. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenyl-amino)-ethylcarbamoyl]-ethyl}-3-fluoro-5-trifluoromethyl-benzamide; * C₂₆H₃₁F₄N₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.62(bs, 1H), 7.75(s, 1H), 7.58(m, 1H), 7.35(m, 4H), 6.88(m, 3H), 4.35(m, 1H), 3.75(s, 3H), 3.56(m, 3H), 3.40(m, 1H), 1.70(m, 7H), 1.37(m, 1H), 1.17(m, 3H), 0.91(m, 2H); LCMS: 510.4(M+H)⁺.

[0266] Example 166. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenyl-amino)-ethylcarbamoyl]-ethyl}-2-fluoro-3-trifluoromethyl-benzamide; * C₂₆H₃₁F₄N₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.50(bs, 1H), 7.87(m, 1H), 7.64(m, 1H), 7.32(m, 2H), 7.15(m, 2H), 6.83(m,

2H), 4.30(m, 1H), 3.72(s, 3H), 3.54(m, 3H), 3.37(m, 1H), 1.65(m, 7H), 1.35(m, 1H), 1.12(m, 3H), 0.89(m, 2H); LCMS: 510.4(M+H)⁺.

[0267] Example 167. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-fluoro-3-methyl-benzamide; * C₂₆H₃₄FN₃O₃; ¹H NMR (CD₃Cl)

5 δ(ppm) 8.62(m, 1H), 7.50(m, 2H), 7.34(m, 2H), 7.09(m, 1H), 6.90(m, 1H): 6.85(m, 2H), 4.32(m, 1H), 3.74(s, 3H), 3.53(m, 3H), 3.38(m, 1H), 2.16(s, 3H), 1.68(m, 7H), 1.34(m, 1H), 1.09(m, 3H) 0.89(m, 2H); LCMS: 456.5(M+H)⁺.

[0268] Example 168. (S)-5-(4-Chloro-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₉H₃₄ClN₃O₄; ¹H NMR

10 (CD₃Cl) δ(ppm) 8.62(m, 1H), 7.53(m, 2H), 7.37(d, J=9.0Hz, 2H), 7.29(m, 2H), 7.19(m, 1H), 6.97(d, J=3.6Hz, 1H), 6.87(m, 2H), 6.58(d, J=3.6Hz, 1H), 4.34(m, 1H), 3.74(s, 3H), 3.60(m, 2H), 3.46(m, 2H), 1.70(m, 7H), 1.39(m, 1H), 1.14(m, 3H), 0.91(m, 2H); LCMS: 524.4(M+H)⁺.

[0269] Example 169. (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-fluoro-4-trifluoromethyl-benzamide; * C₂₆H₃₁F₄N₃O₃; ¹H NMR

15 (CD₃Cl) δ(ppm) 8.57(bs, 1H), 7.81(m, 1H), 7.33(m, 5H), 6.89(m, 2H), 4.32(m, 1H), 3.76(s, 3H), 3.59(m, 3H), 3.43(m, 1H), 1.66(m, 7H), 1.37(m, 1H), 1.15(m, 3H), 0.92(m, 2H); LCMS: 510.5(M+H)⁺.

[0270] Example 170. (S)-4'-Chloro-biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₁H₃₆ClN₃O₃; ¹H NMR (CD₃Cl)

20 δ(ppm) 7.74(bs, 1H), 7.47(d, J=8.4Hz, 2H), 7.40(m, 2H), 7.39(m, 2H), 7.32(m, 4H), 7.13(m, 1H), 6.84(m, 2H), 4.38(m, 1H), 3.73(s, 3H), 3.54(m, 2H), 3.45(m, 1H), 3.36(m, 1H), 1.66(m, 7H), 1.38(m, 1H), 1.12(m, 3H), 0.89(m, 2H); LCMS: 534.4(M+H)⁺.

[0271] Example 171. (S)-3', 5'-Dichloro-biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₁H₃₅Cl₂N₃O₃; ¹H NMR

25 (CD₃Cl) δ(ppm) 8.04(bs, 1H), 7.74(d, J=8.5Hz, 2H), 7.46(d, J=8.4Hz, 2H), 7.34(m, 2H), 7.29(m, 1H), 7.10(m, 2H), 7.00(m, 1H), 6.79(m, 2H), 4.41(m, 1H), 3.71(s, 3H), 3.52(m, 2H), 3.35(m, 2H), 1.64(m, 7H), 1.36(m, 1H), 1.16(m, 3H), 0.89(m, 2H); LCMS: 568.4(³⁵ClM+H)⁺, 570.4(³⁷ClM+H)⁺.

30 **[0272] Example 172.** (S)-3'-Methoxy-biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₂H₃₉N₃O₄; ¹H NMR (CD₃Cl)

δ (ppm) 8.54(bs, 1H), 7.60(d, $J=8.4$ Hz, 2H), 7.41(d, $J=8.4$ Hz, 2H), 7.29(d, $J=9.0$ Hz, 2H), 7.16(m, 1H), 7.08(m, 1H), 6.97(m, 1H), 6.90(m, 1H), 6.76(m, 3H), 4.24(m, 1H), 3.67(s, 3H), 3.64(s, 3H), 3.47(m, 3H), 3.31(m, 1H), 1.57(m, 7H), 1.29(m, 1H), 1.01(m, 3H), 0.81(m, 2H); LCMS: 530.5(M+H)⁺.

5 [0273] **Example 173.** (S)-3'-Chloro-biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₁H₃₆ClN₃O₃; ¹H NMR (CD₃Cl) δ (ppm) 8.66(m, 1H), 7.74(d, $J=8.4$ Hz, 2H), 7.46(m, 3H), 7.39(m, 6H), 6.85(m, 2H), 4.37(m, 1H), 3.73(s, 3H), 3.51(m, 3H), 3.39(m, 1H), 1.69(m, 7H), 1.38(m, 1H), 1.13(m, 3H), 0.90(m, 2H); LCMS: 534.5(M+H)⁺.

10 [0274] **Example 174.** (S)-2'-Methoxy-biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₂H₃₉N₃O₄; ¹H NMR (CD₃Cl) δ (ppm) 8.66(m, 1H), 7.69(d, $J=8.4$ Hz, 2H), 7.48(d, $J=8.4$ Hz, 2H), 7.39(d, $J=9.0$ Hz, 2H), 7.27(m, 1H), 7.20(m, 1H), 7.08(m, 1H), 6.89(m, 4H), 4.34(m, 1H), 3.73(s, 3H), 3.72(s, 3H), 3.56(m, 3H), 3.39(m, 1H), 1.67(m, 7H), 1.38(m, 1H), 1.13(m, 3H), 0.90(m, 2H); LCMS: 530.5(M+H)⁺.

15 [0275] **Example 175.** (S)-4'-Chloro-biphenyl-3-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₁H₃₆ClN₃O₃; ¹H NMR (CD₃Cl) δ (ppm) 8.43(m, 1H), 7.49(m, 1H), 7.42(m, 1H), 7.31(m, 8H), 6.84(d, $J=9.0$ Hz, 2H), 5.98(m, 1H), 3.96(m, 1H), 3.74(s, 3H), 3.59(m, 2H), 3.45(m, 1H), 3.38(m, 1H), 1.58(m, 3H), 1.46(m, 3H), 1.09(m, 4H), 0.76(m, 3H); LCMS: 534.4(M+H)⁺.

20 [0276] **Example 176.** (S)-4-Benzo[1,3]dioxol-5-yl-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide; * C₃₂H₃₇N₃O₅; ¹H NMR (CD₃Cl) δ (ppm) 8.68(m, 1H), 7.68(d, $J=8.4$ Hz, 2H), 7.40(m, 4H), 7.11(m, 1H), 6.97(m, 2H), 6.88(m, 2H), 6.79(m, 1H), 5.93(s, 2H), 4.33(m, 1H), 3.74(s, 3H), 3.55(m, 3H), 3.40(m, 1H), 1.70(m, 7H), 1.39(m, 1H), 1.13(m, 3H) 0.92(m, 2H); LCMS: 544.5(M+H)⁺.

25 [0277] **Example 177.** (S)-5-Bromo-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₃H₃₀BrN₃O₄; ¹H NMR (CD₃Cl) δ (ppm) 8.3(m, 1H), 7.21(m, 2H), 6.83(m, 1H), 6.72(m, 3H), 6.20(d, $J=3.5$ Hz, 1H), 4.14(dd, $J=3.8$ Hz, $J=5.2$ Hz, 1H), 3.61(s, 3H), 3.45(m, 2H), 3.34(m, 1H), 3.28(m, 1H), 1.52(m, 7H), 1.22(m, 1H), 1.02(m, 3H), 0.77(m, 2H); LCMS: 492.3(⁷⁹BrM+H)⁺, 494.3(⁸⁰BrM+H)⁺.

[0278] **Example 178.** (S)-3,5-Dibromo-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide; $^* \text{C}_{25}\text{H}_{31}\text{Br}_2\text{N}_3\text{O}_3$; ^1H NMR (CD_3Cl) δ (ppm) 8.61(m, 1H), 7.78(m, 2H), 7.69(s, 1H), 7.45(m, 1H), 7.31(m, 2H), 6.86(m, 2H), 4.36(m, 1H), 3.74(s, 3H), 3.59(m, 1H), 3.42(m, 3H), 1.60(m, 7H), 1.35(m, 1H), 1.14(m, 3H), 0.86(m, 2H); LCMS:

580.2, 582.2, 584.3.

[0279] **Example 179.** (S)-3,5-Dichloro-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide; $^* \text{C}_{25}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_3$; ^1H NMR (CD_3Cl) δ (ppm) 8.60(m, 1H), 7.58(s, 2H), 7.51(m, 1H), 7.38(m, 1H), 7.30(d, $J=9.0\text{Hz}$, 2H), 6.83(m, 2H), 4.37(m, 1H), 3.74(s, 3H), 3.57(m, 1H), 3.43(m, 3H), 1.61(m, 7H), 1.32(m, 1H), 1.15(m, 3H), 0.85(m, 2H); LCMS: 492.4($^{35}\text{ClM}+\text{H}$) $^+$, 494.3($^{37}\text{ClM}+\text{H}$) $^+$.

[0280] **Example 180.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3,5-dimethoxy-benzamide; $^* \text{C}_{27}\text{H}_{37}\text{N}_3\text{O}_5$; ^1H NMR (CD_3Cl) δ (ppm) 8.59(m, 1H), 7.37(m, 2H), 7.11(m, 1H), 6.85(m, 2H), 6.76(m, 2H), 6.49(m, 1H), 4.30(m, 1H), 3.73(s, 3H), 3.65(s, 6H), 3.56(m, 2H), 3.44(m, 2H), 1.66(m, 7H), 1.35(m, 1H), 1.15(m, 3H), 0.89(m, 2H); LCMS: 484.5(M+H) $^+$.

[0281] **Example 181.** (S)-Biphenyl-3-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; $^* \text{C}_{31}\text{H}_{37}\text{N}_3\text{O}_3$; ^1H NMR (CD_3Cl) δ (ppm) 8.71(m, 1H), 7.90(s, 1H), 7.71(m, 1H), 7.65(m, 1H), 7.44(m, 9H), 6.89(d, $J=9.0\text{Hz}$, 2H), 4.40(m, 1H), 3.77(s, 3H), 3.63(m, 3H), 3.50(m, 1H), 1.72(m, 7H), 1.37(m, 1H), 1.19(m, 3H), 0.96(m, 2H); LCMS: 500.5(M+H) $^+$.

[0282] **Example 182.** (S)-5-Bromo-thiophene-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; $^* \text{C}_{23}\text{H}_{30}\text{BrN}_3\text{O}_3\text{S}$; ^1H NMR (CD_3Cl) δ (ppm) 8.63(m, 1H), 7.47(d, $J=5.5\text{Hz}$, 1H), 7.35(m, 2H), 7.29(s, $J=4.0\text{Hz}$, 1H), 6.93(s, $J=4.0\text{Hz}$, 1H), 6.88(m, 2H), 4.29(m, 1H), 3.76(s, 3H), 3.57(m, 1H), 3.47(m, 2H), 3.40(m, 1H), 1.60(m, 7H), 1.34(m, 1H), 1.11(m, 3H), 0.84(m, 2H); LCMS: 508.3($^{79}\text{BrM}+\text{H}$) $^+$, 510.3($^{80}\text{BrM}+\text{H}$) $^+$.

[0283] **Example 183.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-phenoxy-benzamide; $^* \text{C}_{31}\text{H}_{37}\text{N}_3\text{O}_4$; ^1H NMR (CD_3Cl) δ (ppm) 8.62(m, 1H), 7.60(d, $J=8.8\text{Hz}$, 2H), 7.38(d, $J=9.0\text{Hz}$, 2H), 7.28(m, 2H), 7.09(m, 2H), 6.94(m, 2H), 6.86(m, 4H), 4.30(m, 1H), 3.74(s, 3H), 3.54(m, 3H), 3.37(m, 1H), 1.66(m, 7H), 1.37(m, 1H), 1.12(m, 3H), 0.87(m, 2H); LCMS: 516.5(M+H) $^+$.

[0284] **Example 184.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-phenoxy-benzamide; * C₃₁H₃₇N₃O₄; ¹H NMR (CD₃Cl) δ(ppm) 8.60(m, 1H), 7.34(m, 3H), 7.23(m, 4H), 7.02(m, 3H), 6.87(m, 2H), 6.81(d, J=9Hz, 2H), 4.28(m, 1H), 3.73(s, 3H), 3.52(m, 3H), 3.38(m, 1H), 1.64(m, 7H), 1.34(m, 1H), 1.10(m, 3H), 0.88(m, 2H); LCMS: 516.5(M+H)⁺.

[0285] **Example 185.** (S)-1H-Indole-3-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₇H₃₄N₄O₃; LCMS: 463.5(M+H)⁺.

[0286] **Example 186.** (S)-Benzothiazole-6-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₆H₃₂N₄O₃S; ¹H NMR (CD₃Cl) δ(ppm) 9.07(bs, 1H), 8.62(bs, 1H), 8.33(s, 1H), 8.05(m, 1H), 7.79(m, 1H), 7.37(m, 3H), 6.87(m, 2H), 4.40(dd, J=5.2Hz, J=9.3Hz, 1H), 3.74(s, 3H), 3.55(m, 3H), 3.41(m, 1H), 1.69(m, 7H), 1.41(m, 1H), 1.16(m, 3H), 0.89(m, 2H); LCMS: 481.4(M+H)⁺.

[0287] **Example 187.** (S)-2-Amino-benzothiazole-6-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₆H₃₃N₅O₃S; ¹H NMR (CD₃Cl) δ(ppm) 9.65(bs, 1H), 8.17(m, 1H), 7.71(s, 1H), 7.57(d, J=8.6Hz, 2H), 7.48(d, J=9.0Hz, 2H), 7.29(m, 2H), 6.92(d, J=9.0Hz, 2H), 4.25(m, 1H), 3.76(s, 3H), 3.62(m, 3H), 3.43(m, 1H), 1.73(m, 8H), 1.18(m, 3H), 0.93(m, 2H); LCMS: 496.4(M+H)⁺.

[0288] **Example 188.** (S)-4-Methyl-2-(4-trifluoromethyl-phenyl)-thiazole-5-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₀H₃₅F₃N₄O₃S; ¹H NMR (CD₃Cl) δ(ppm) 8.61(m, 1H), 7.91(d, J=8.1Hz, 2H), 7.61(d, J=8.3Hz, 2H), 7.38(m, 2H), 6.90(m, 2H), 6.72(d, J=4.8Hz, 1H), 4.31(m, 1H), 3.75(s, 3H), 3.61(m, 3H), 3.44(m, 1H), 2.54(s, 3H), 1.67(m, 7H), 1.37(m, 1H), 1.17(m, 3H), 0.92(m, 2H); LCMS: 589.4(M+H)⁺.

[0289] **Example 189.** (S)-4-(4-Chloro-phenyl)-thiophene-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₉H₃₄ClN₃O₃S; ¹H NMR (CD₃Cl) δ(ppm) 8.72(m, 1H), 7.83(m, 1H), 7.46(s, 1H), 7.36(m, 4H), 7.23(m, 2H), 6.86(d, J=9.0Hz, 2H), 4.38(m, 1H), 3.74(s, 3H), 3.55(m, 1H), 3.44(m, 3H), 1.60(m, 7H), 1.39(m, 1H), 1.10(m, 3H), 0.89(m, 2H); LCMS: 540.4(M+H)⁺.

[0290] **Example 190.** (S)-2-Methyl-5-trifluoromethyl-oxazole-4-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₄H₃₁F₃N₄O₄; ¹H NMR (CD₃Cl) δ(ppm) 8.62(m, 1H), 7.68(m, 1H), 7.40(m, 2H), 6.92(m, 2H), 4.26(m, 1H),

3.81(s, 3H), 3.74(m, 2H), 3.59(m, 1H), 3.35(m, 1H), 2.51(s, 3H), 1.74(m, 7H), 1.44(m, 1H), 1.20(m, 3H), 0.97(m, 2H); LCMS: 497.4(M+H)⁺.

[0291] **Example 191.** (S)-4-(4-Methoxy-phenyl)-thiophene-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₀H₃₇N₃O₄S;

5 ¹H NMR (CD₃Cl) δ(ppm) 8.72(m, 1H), 7.82(m, 1H), 7.74(m, 1H), 7.35(m, 5H), 6.85(m, 2H), 6.78(m, 2H), 4.37(m, 1H), 3.73(s, 3H), 3.69(s, 3H), 3.53(m, 1H), 3.42(m, 3H), 1.66(m, 7H), 1.41(m, 1H), 1.10(m, 3H), 0.86(m, 2H); LCMS: 536.5(M+H)⁺.

[0292] **Example 192.** N-{2-Cyclohexyl-1-(S)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-2-(S)-phenyl-propionamide; * C₂₇H₃₇N₃O₃; ¹H NMR (CD₃Cl) δ(ppm)

10 8.30(m, 1H), 7.23(m, 6H), 7.18(m, 1H), 6.83(m, 2H), 6.26(m, 1H), 3.96(m, 1H), 3.74(s, 3H), 3.56(m, 2H), 3.49(m, 1H), 3.38(m, 1H), 3.31(m, 1H), 1.49(m, 7H), 1.39(d, J=7.2Hz, 3H), 1.00(m, 4H), 0.78(m, 2H); LCMS: 452.5(M+H)⁺.

[0293] **Example 193.** (S)-5-(2-Chloro-5-trifluoromethyl-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; *

15 C₃₀H₃₃ClF₃N₃O₄; ¹H NMR (CD₃Cl) δ(ppm) 8.56(m, 1H), 7.97(m, 1H), 7.50(m, 1H), 7.40(m, 1H), 7.39(m, 2H), 7.25(m, 1H), 7.10(d, J=3.7, 1H), 7.01(d, J=3.7, 1H), 6.89(m, 2H), 4.29(m, 1H), 3.75(s, 3H), 3.59(m, 3H), 3.46(m, 1H), 1.72(m, 7H), 1.40(m, 1H), 1.13(m, 3H), 0.91(m, 2H); LCMS: 592.4(M+H)⁺.

[0294] **Example 194.** (S)-2'-Chloro-biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-

20 trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₁H₃₃ClF₃N₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 7.72(d, J=8.4Hz, 2H), 7.52(m, 1H), 7.42(m, 3H), 7.24(m, 3H), 7.04(m, 2H), 6.92(m, 2H), 6.83(m, 1H), 4.47(m, 1H), 3.49(m, 2H), 3.33(m, 2H), 1.64(m, 7H), 1.34(m, 1H), 1.13(m, 3H), 0.91(m, 2H); LCMS: 558.4(M+H)⁺.

[0295] **Example 195.** (S)-1-(5-Bromo-pyrimidin-2-yl)-piperidine-4-carboxylic acid {2-

25 cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₈H₃₉BrN₆O₃; ¹H NMR (CD₃Cl) δ(ppm) 8.51(m, 1H), 8.17(m, 2H), 7.33(m, 2H), 6.84(m, 2H), 6.44(d, J=4.7Hz, 1H), 4.60(m, 2H), 4.07(m, 1H), 3.74(s, 3H), 3.53(m, 3H), 3.31(m, 1H), 2.76(m, 2H), 2.43(m, 1H), 1.81(m, 1H), 1.55(m, 10H), 1.27(m, 1H), 1.10(m, 3H), 0.86(m, 2H); LCMS: 587.4(⁷⁹BrM+H)⁺, 589.5(⁸⁰BrM+H)⁺.

[0296] **Example 196.** (S)-N-{2-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-

ethylcarbamoyl]-ethyl}-4-(4,6-dimethyl-pyrimidin-2-ylamino)-benzamide; * C₃₁H₄₀N₆O₃; ¹H

NMR (CD₃Cl) δ (ppm) 12.17(bs, 1H), 8.65(m, 1H), 7.68(m, 4H), 7.37(m, 3H), 6.85(m, 2H), 6.57(s, 1H), 4.35(m, 1H), 3.74(s, 3H), 3.55(m, 2H), 3.41(m, 2H), 2.49(s, 6H), 1.66(m, 7H), 1.37(m, 1H), 1.14(m, 3H), 0.88(m, 2H); LCMS: 545.5(M+H)⁺.

[0297] **Example 197.** (S)-1-(5-Bromo-pyrimidin-2-yl)-piperidine-3-carboxylic acid {2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₈H₃₉BrN₆O₃; ¹H NMR (CD₃Cl) δ (ppm) 8.42(m, 1H), 8.32(s, 1H), 8.20(s, 1H), 7.36(m, 2H), 7.03(m, 1H), 6.87(m, 2H), 3.90(m, 2H), 3.74(s, 3H), 3.59(m, 5H), 2.45(m, 1H), 1.80(m, 2H), 1.54(m, 1H), 1.03(m, 6H); LCMS: 587.5(⁷⁹BrM+H)⁺, 589.4(⁸⁰BrM+H)⁺.

[0298] **Example 198.** (S)-3'-Fluoro-biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₁H₃₃F₄N₃O₃; ¹H NMR (CD₃Cl) δ (ppm) 7.73(m, 2H), 7.53(d, *J*=8.4Hz, 2H), 7.52(m, 3H), 7.20(m, 1H), 6.98(m, 3H), 6.71(m, 3H), 4.52(m, 1H), 3.46(m, 2H), 3.26(m, 2H), 1.63(m, 7H), 1.31(m, 1H), 1.09(m, 3H), 0.90(m, 2H); LCMS: 572.5(M+H)⁺.

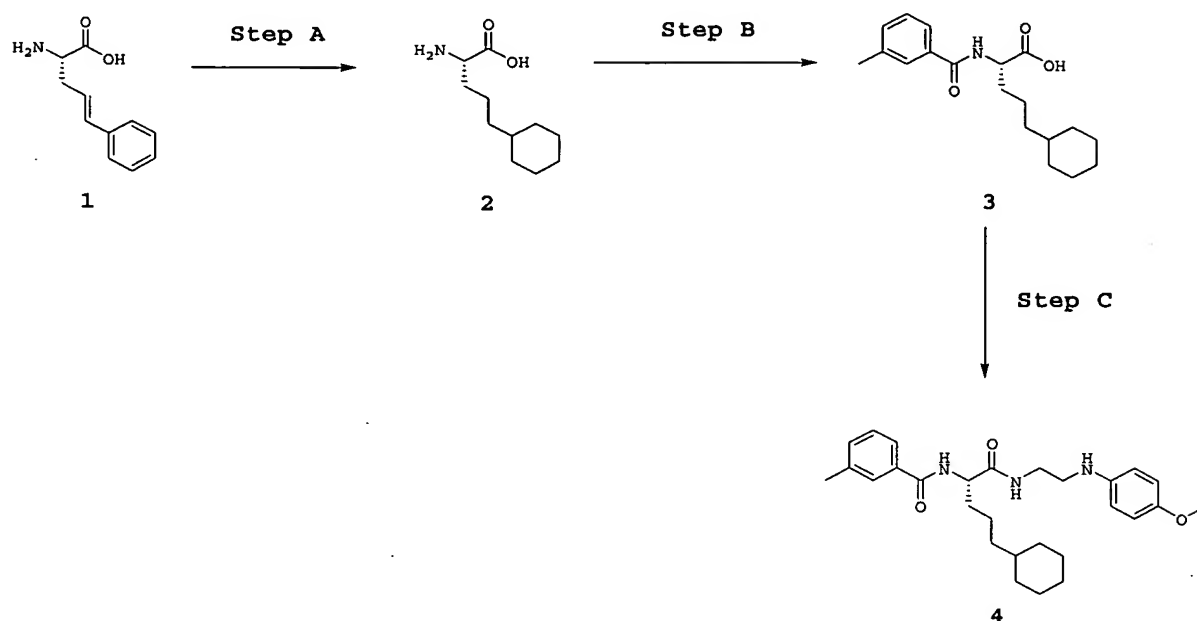
[0299] **Example 199.** (S)-3-Aminomethyl-N-{2-cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide; * C₂₆H₃₆N₄O₃; ¹H NMR (CD₃Cl) δ (ppm) 8.44(bs, 3H), 7.91(s, 1H), 7.61(m, 1H), 7.33(m, 4H), 6.86(m, 2H), 4.42(m, 1H), 3.98(bs, 2H), 3.74(s, 3H), 3.37(m, 4H), 2.59(s, 2H), 1.68(m, 7H), 1.35(m, 1H), 1.14(m, 3H), 0.89(m, 2H); LCMS: 453.5(M+H)⁺.

[0300] **Example 200.** (S)-N-{2-Cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-morpholin-4-ylmethyl-benzamide; * C₃₀H₃₉F₃N₄O₄; LCMS: 557.5(M+H)⁺.

[0301] **Example 201.** (S)-5-(2-Fluoro-phenyl)-thiophene-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₉H₃₁F₄N₃O₃S; ¹H NMR (CD₃Cl) δ (ppm) 7.51(m, 1H), 7.45(m, 1H), 7.39(m, 1H), 7.31(m, 1H), 7.23(m, 1H), 7.08(m, 2H), 6.99(m, 2H), 6.77(m, 3H), 4.48(dd, *J*=6.8Hz, *J*=15.2Hz, 1H), 3.47(m, 2H), 3.26(m, 2H), 1.62(m, 7H), 1.32(m, 1H), 1.10(m, 3H), 0.88(m, 2H); LCMS: 578.4(M+H)⁺.

[0302] **Example 202.** (S)-N-{3-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-homo-cyclohexyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (12 mg, 53%): MS calcd. for C₂₇H₃₈N₃O₃ (M+H)⁺ 452.29, found 452.3.

[0303] **Example 203.** (S)-N-{4-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-butyl}-3-methyl-benzamide.



Step A: L-Styryl alanine **1** (50 mg, 0.26 mmol) was dissolved in MeOH (5 mL) and placed into a Parr-hydrogenation apparatus. Catalytic amounts of rhodium (5% on Al₂O₃) were added and the reaction vessel was placed under a hydrogen at 55 psi. The mixture was shaken for 20 h at room temperature, then filtered over celite. The organic solvent was removed in vacuo to yield (S)-2-Amino-5-cyclohexyl-pentanoic acid **2** (52 mg, quant.) as a white solid: ¹H-NMR (400MHz, CD₃OD) δ = 3.50 (dd, *J* = 5.2, *J* = 6.9, 1H), 1.88-1.64 (m, 7H), 1.47-1.37 (m, 2H), 1.31-1.11 (m, 6H), 0.94-0.86 (m, 2H). MS calcd. for C₁₁H₂₂NO₂ (M+H⁺) 200.17, found 200.4.

Step B: (S)-2-Amino-5-cyclohexyl-pentanoic acid **2** (24 mg, 0.12 mmol) was dissolved in H₂O (1 mL) containing equimolar amounts of NaOH (5 mg, 0.12 mmol). The solution was cooled to 0 °C, then m-Toluic acid chloride (16 μL, 0.12 mmol) was added dropwise under vigorous stirring. The mixture was allowed to warm to room temperature and stirred for approx. 12 h. After acidification with 1 M HCl (1 mL), the product was extracted from the reaction mixture with DCM (4 mL). The organic layer was separated and the solvent was removed in vacuo to yield (S)-5-Cyclohexyl-2-(3-methyl-benzoylamino)-pentanoic acid **3** (19 mg, 50%) as a white solid: ¹H-NMR (400MHz, CD₃OD) δ = 7.67-7.62 (m, 2H), 7.37-7.31 (m, 2H), 4.56 (dd, *J* = 5.0, *J* = 9.4, 1H), 2.40 (s, 3H), 1.96-0.84 (m, 17H). MS calcd. for C₁₉H₂₈NO₃ (M+H⁺) 318.21, found 318.4.

Step C: (S)-5-Cyclohexyl-2-(3-methyl-benzoylamino)-pentanoic acid **3** (19 mg, 0.06 mmol) was dissolved in DCM (2 mL), HOBT (20 mg, 0.14 mmol) and DIC (23 μ L, 0.14 mmol) were added and the solution was stirred for 10 min at room temperature. N-(4-Methoxyphenyl)-ethane-1,2-diamine (24 mg, 0.14 mmol) was added and the solution was stirred for 3 h at room temperature. The solvent was removed in vacuo, and the remainder was purified by reverse HPLC to afford the title compound (S)-N-{4-Cyclohexyl-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-butyl}-3-methyl-benzamide **4** (15 mg, 0.03 mmol, 50%) as a white solid: $^1\text{H-NMR}$ (400MHz, CD_3OD) δ = 7.70-7.65 (m, 2H), 7.38-7.32 (m, 4H), 7.03-6.99 (m, 2H), 4.39 (dd, J = 6.4, J = 8.4, 1H), 3.80 (s, 3H), 3.67-3.61 (m, 1H), 3.54-3.41 (m, 3H), 2.38 (s, 3H), 1.88-1.80 (m, 2H), 1.72-1.66 (m, 5H), 1.64-1.36 (m, 2H), 1.29-1.07 (m, 6H), 0.92-0.83 (m, 2H). MS calcd. for $\text{C}_{28}\text{H}_{40}\text{N}_3\text{O}_3$ ($\text{M}+\text{H}^+$) 466.31, found 466.5.

[0304] Example 204. (S)-N-{[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-phenyl-methyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-phenylglycine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (7 mg, 34%): MS calcd. for $\text{C}_{25}\text{H}_{28}\text{N}_3\text{O}_3$ ($\text{M}+\text{H}^+$) 418.21, found 418.2.

[0305] Example 205. (S)-5-(4-Trifluoromethyl-phenyl)-thiophene-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * $\text{C}_{30}\text{H}_{31}\text{F}_6\text{N}_3\text{O}_3\text{S}$; $^1\text{H NMR}$ (CD_3Cl) δ (ppm) 7.58(m, 4H), 7.49(m, 1H), 7.44(d, J =4Hz, 1H), 7.24(d, J =4Hz, 1H), 7.07(d, J =8.8Hz, 2H), 6.96(d, J =8.8Hz, 2H), 6.78(m, 1H), 4.42(dd, J =6.4Hz, J =15.2Hz, 1H), 3.53(m, 2H), 3.32(m, 2H), 1.63(m, 7H), 1.31(m, 1H), 1.14(m, 3H), 0.90(m, 2H); LCMS: 628.4($\text{M}+\text{H}$) $^+$.

[0306] Example 206. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-3-phenyl-propyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-homo phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (9 mg, 40%): MS calcd. for $\text{C}_{27}\text{H}_{32}\text{N}_3\text{O}_3$ ($\text{M}+\text{H}^+$) 446.24, found 446.2.

[0307] Example 207. (S)-5-(4-Trifluoromethoxy-phenyl)-thiophene-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * $\text{C}_{30}\text{H}_{31}\text{F}_6\text{N}_3\text{O}_4\text{S}$; $^1\text{H NMR}$ (CD_3Cl) δ (ppm) 7.49(m, 3H), 7.43(d, J =4Hz, 1H), 7.17(m, 3H), 7.06(m, 2H), 6.92(m, 2H), 6.83(m, 1H), 4.43(m, 1H), 3.51(m, 2H), 3.25(m, 2H), 1.62(m, 7H), 1.32(m, 1H), 1.14(m, 3H), 0.89(m, 2H); LCMS: 644.4($\text{M}+\text{H}$) $^+$.

[0308] **Example 208.** (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-4-phenyl-but-3-enyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-styryl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (9
5 mg, 39%): MS calcd. for $C_{28}H_{32}N_3O_3$ ($M+H^+$) 458.24, found 458.2.

[0309] **Example 209.** (S)-5-(3-Trifluoromethoxy-phenyl)-thiophene-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; *
 $C_{30}H_{31}F_6N_3O_4S$; 1H NMR (CD_3Cl) δ (ppm) 7.41(m, 2H), 7.34(m, 2H), 7.20(m, 1H), 7.16(m, 2H), 6.99(d, $J=8.0Hz$, 2H), 6.74(m, 3H), 4.47(m, 1H), 3.47(m, 2H), 3.26(m, 2H), 1.61(m,
10 7H), 1.30(m, 1H), 1.12(m, 3H), 0.88(m, 2H); LCMS: 644.4($M+H$) $^+$.

[0310] **Example 210.** (S)-5-(2-Methoxy-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * $C_{30}H_{34}F_3N_3O_5$; 1H NMR (CD_3Cl) δ (ppm) 7.77(d, 1H), 7.29(m, 2H), 7.07(d, $J=3.6Hz$, 1H), 6.98(m, 5H), 6.84(m, 2H), 6.77(m, 1H), 4.46(m, 1H), 3.87(s, 3H), 3.49(m, 2H), 3.30(m, 2H), 1.63(m, 7H),
15 1.32(m, 1H), 1.14(m, 3H), 0.90(m, 2H); LCMS: 628.4($M+H$) $^+$.

[0311] **Example 211.** N-{(4-Methoxy-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide. ** Following the procedure of Example 227, except substituting 4-fluoro-phenyl glycine for 4-methoxy-phenyl glycine in Step A, the title compound was prepared as a white solid (16 mg, 30%): 1H -NMR (400MHz, CD_3OD) δ =
20 7.69-6.94 (m, 12H), 5.43 (s, 1H), 3.82 (s, 3H), 3.80 (s, 3H), 3.71-3.37 (m, 4H), 2.37 (s, 3H). MS calcd. for $C_{26}H_{30}N_3O_4$ ($M+H^+$) 448.22, found 448.5.

[0312] **Example 212.** (S)-5-(2-Fluoro-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * $C_{29}H_{31}F_4N_3O_4$; 1H NMR (CD_3Cl) δ (ppm) 7.75(m, 2H), 7.27(m, 1H), 7.12(m, 7H), 6.99(m, 1H), 6.83(m, 1H),
25 4.42(dd, $J=6.4Hz$, $J=14.8Hz$, 1H), 3.55(m, 2H), 3.38(m, 2H), 1.65(m, 7H), 1.35(m, 1H), 1.13(m, 3H), 0.91(m, 2H); LCMS: 644.4($M+H$) $^+$.

[0313] **Example 213.** (S)-5-(4-Trifluoromethyl-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; *
 $C_{30}H_{31}F_6N_3O_4$; 1H NMR (CD_3Cl) δ (ppm) 7.72(d, $J=8.0$, 2H); 7.59(m, 3H); 7.06(m, 6H);
30 6.75(d, $J=3.6Hz$, 1H); 4.45(m, 1H); 3.54(m, 2H); 3.36(m, 2H); 1.65(m, 7H); 1.33(m, 1H); 1.13(m, 3H); 0.89(m, 2H); LCMS: 574.5($M+H$) $^+$.

[0314] **Example 214.** N-{{(2-Benzyloxy-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide. ** Following the procedure of Example 227, except substituting 4-fluoro-phenyl glycine for 2-benzyloxy-phenyl glycine in Step A, the title compound was prepared as a white solid (7 mg, 11%): ¹H-NMR (400MHz, CD₃OD) δ =
5 7.64-6.99 (m, 17H), 5.94 (s, 1H), 5.18 (s, 2H), 3.82 (s, 3H), 3.63-3.37 (m, 4H), 2.36 (s, 3H). MS calcd. for C₃₂H₃₄N₃O₄ (M+H⁺) 524.25, found 524.6.

[0315] **Example 215.** (S)-5-(4-Trifluoromethoxy-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; *
C₃₀H₃₁F₆N₃O₅; ¹H NMR (CD₃Cl) δ (ppm) 7.65(d, J=8.8, 2H), 7.56(m, 1H), 7.21(m, 1H),
10 7.04(m, 6H), 6.64(d, J=3.6Hz, 1H), 4.42(dd, J=2.4Hz, J=6.4Hz, 1H), 3.54(m, 2H), 3.33(m, 2H), 1.66(m, 7H), 1.33(m, 1H), 1.15(m, 3H), 0.91(m, 2H); LCMS: 562.4(M+H)⁺.

[0316] **Example 216.** N-{{(2-Chloro-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide. ** Following the procedure of Example 227, except substituting 4-fluoro-phenyl glycine for 2-chloro-phenyl glycine in Step A, the title
15 compound was prepared as a white solid (20 mg, 37%): ¹H-NMR (400MHz, CD₃OD) δ = 7.70-7.01 (m, 12H), 5.98 (s, 1H), 3.82 (s, 3H), 3.75-3.43 (m, 4H), 2.37 (s, 3H). MS calcd. for C₂₅H₂₇ClN₃O₃ (M+H⁺) 452.17, found 452.5.

[0317] **Example 217.** N-{{(4-Benzyloxy-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide. ** Following the procedure of Example 227, except substituting 4-fluoro-phenyl glycine for 4-benzyloxy-phenyl glycine in Step A, the
20 title compound was prepared as a white solid (14 mg, 22%): ¹H-NMR (400MHz, CD₃OD) δ = 7.69-7.01 (m, 17H), 5.42 (s, 1H), 5.10 (s, 2H), 3.81 (s, 3H), 3.72-3.35 (m, 4H), 2.37 (s, 3H). MS calcd. for C₃₂H₃₄N₃O₄ (M+H⁺) 524.25, found 524.5.

[0318] **Example 218.** N-{{[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-naphthalen-1-yl-methyl}-3-methyl-benzamide. ** Following the procedure of Example 227, except
25 substituting 4-fluoro-phenyl glycine for 1-naphtyl glycine in Step A, the title compound was prepared as a white solid (20 mg, 36%): ¹H-NMR (400MHz, CD₃OD) δ = 8.15-7.04 (m, 15H), 6.39 (s, 1H), 3.82 (s, 3H), 3.80-3.42 (m, 4H), 2.34 (s, 3H). MS calcd. for C₂₉H₃₀N₃O₃ (M+H⁺) 468.23, found 468.5.

[0319] **Example 219.** (S)-5-(2-Trifluoromethyl-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; *
30 C₃₀H₃₁F₆N₃O₄; ¹H NMR (CD₃Cl) δ(ppm) 7.83(d, J=8.0, 1H), 7.70(m, 1H), 7.64(m, 1H),

7.30(m, 1H), 7.25(m, 1H), 7.13(m, 3H), 6.91(m, 2H), 6.74(m, 2H), 4.47(dd, $J=3.2\text{Hz}$, $J=6.0\text{Hz}$, 1H), 3.58(m, 2H), 3.37(m, 2H), 1.69(m, 7H), 1.40(m, 1H), 1.21(m, 3H), 0.98(m, 2H); LCMS: 612.4(M+H)⁺.

[0320] Example 220. N-{{2-(4-Methoxy-phenylamino)-ethylcarbamoyl}-o-tolyl-methyl}-3-methyl-benzamide. ** Following the procedure of Example 227, except substituting 4-fluoro-phenyl glycine for 2-methyl-phenyl glycine in Step A, the title compound was prepared as a white solid (19 mg, 37%): ¹H-NMR (400MHz, CD₃OD) δ = 7.70-7.05 (m, 12H), 5.78 (s, 1H), 3.83 (s, 3H), 3.77-3.41 (m, 4H), 2.45 (s, 3H), 2.37 (s, 3H). MS calcd. for C₂₆H₃₀N₃O₃ (M+H⁺) 432.23, found 432.23.

[0321] Example 221. (S)-N-{2-Cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-4-[1,2,4]triazol-1-yl-benzamide; * C₂₇H₃₁F₃N₆O₃; ¹H NMR (CD₃Cl) δ (ppm) 8.60(s, 1H), 8.07(s, 1H), 7.82(m, 2H), 7.68(d, $J=8.8\text{Hz}$, 2H), 7.32(m, 1H), 7.04(d, $J=8.4\text{Hz}$, 2H), 6.92(d, $J=6.8\text{Hz}$, 1H), 6.84(m, 2H), 4.49(m, 1H), 3.50(m, 2H), 3.30(m, 2H), 1.65(m, 7H), 1.32(m, 1H), 1.10(m, 3H), 0.88(m, 2H); LCMS: 545.4(M+H)⁺.

[0322] Example 222. N-{{(2,4-Dichloro-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide. ** Following the procedure of Example 227, except substituting 4-fluoro-phenyl glycine for 2,4-dichloro-phenyl glycine in Step A, the title compound was prepared as a white solid (10 mg, 17%): ¹H-NMR (400MHz, CD₃OD) δ = 7.70-7.04 (m, 11H), 5.93 (s, 1H), 3.82 (s, 3H), 3.74-3.44 (m, 4H), 2.38 (s, 3H). MS calcd. for C₂₅H₂₆Cl₂N₃O₃ (M+H⁺) 486.14, found 486.4.

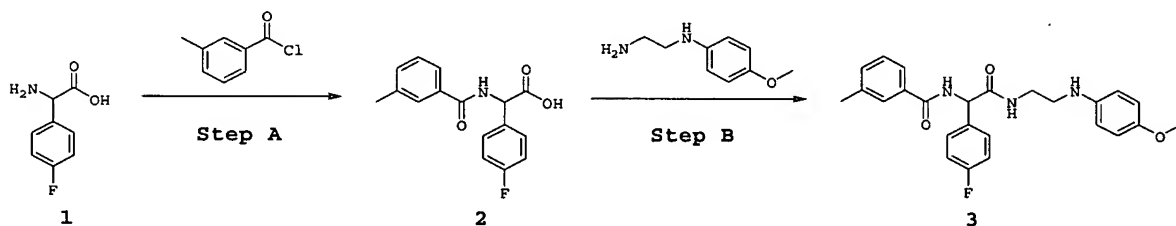
[0323] Example 223. N-{{(2,3-Dichloro-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide. ** Following the procedure of Example 227, except substituting 4-fluoro-phenyl glycine for 2,3-dichloro-phenyl glycine in Step A, the title compound was prepared as a white solid (32 mg, 55%): ¹H-NMR (400MHz, CD₃OD) δ = 7.70-7.00 (m, 11H), 6.03 (s, 1H), 3.80 (s, 3H), 3.70-3.43 (m, 4H), 2.37 (s, 3H). MS calcd. for C₂₅H₂₆Cl₂N₃O₃ (M+H⁺) 486.14, found 486.4.

[0324] Example 224. N-{{(2,4-Dimethyl-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide. ** Following the procedure of Example 227, except substituting 4-fluoro-phenyl glycine for 2,4-dimethyl-phenyl glycine in Step A, the title compound was prepared as a white solid (16 mg, 30%): ¹H-NMR (400MHz, CD₃OD) δ = 7.69-7.01 (m, 11H), 5.73 (s, 1H), 3.82 (s, 3H), 3.75-3.40 (m, 4H), 2.40 (s, 3H), 2.37 (s, 3H), 2.30 (s, 3H). MS calcd. for C₂₇H₃₂N₃O₃ (M+H⁺) 446.24, found 446.5.

[0325] Example 225. N-[(2,4-Dimethoxy-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl]-3-methyl-benzamide. ** Following the procedure of Example 227, except substituting 4-fluoro-phenyl glycine for 2,4-dimethoxy-phenyl glycine in Step A, the title compound was prepared as a white solid (16 mg, 28%): ¹H-NMR (400MHz, CD₃OD) δ = 7.68-6.54 (m, 11H), 5.78 (s, 1H), 3.87 (s, 3H), 3.82 (s, 3H), 3.80 (s, 3H), 3.71-3.40 (m, 4H), 2.37 (s, 3H). MS calcd. for C₂₇H₃₂N₃O₅ (M+H⁺) 478.23, found 478.5.

[0326] Example 226. N-[[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-thiophen-2-yl-methyl]-3-methyl-benzamide. ** Following the procedure of Example 227, except substituting 4-fluoro-phenyl glycine for 2-thienyl glycine in Step A, the title compound was prepared as a white solid (5 mg, 10%): ¹H-NMR (400MHz, CD₃OD) δ = 7.70-7.04 (m, 11H), 5.80 (s, 1H), 3.82 (s, 3H), 3.75-3.40 (m, 4H), 2.38 (s, 3H). MS calcd. for C₂₃H₂₆N₃O₃S (M+H⁺) 424.17, found 424.5.

[0327] Example 227. N-[(4-Fluoro-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl]-3-methyl-benzamide.



Step A: 4-Fluoro-phenyl glycine **1** (20 mg, 0.12 mmol) was dissolved in H₂O (1 mL) containing equimolar amounts of NaOH (5 mg, 0.12 mmol). The solution was cooled to 0 °C, then m-toluyoyl chloride (16 μL, 0.12 mmol) was added dropwise under vigorous stirring. The mixture was allowed to warm to room temperature and stirred for approx. 12 h. After acidification with 1 M HCl (1 mL), the product was extracted from the reaction mixture with DCM (4 mL). The organic layer was separated and the solvent was removed in vacuo to yield (4-Fluoro-phenyl)-(3-methyl-benzoylamino)-acetic acid **2** (34 mg, 0.12 mmol, quant.) as a white solid: ¹H-NMR (400MHz, CD₃OD) δ = 7.67-7.08 (m, 8H), 5.66 (s, 1H), 2.38 (s, 3H). MS calcd. for C₁₆H₁₅FO₃ (M+H⁺) 288.10, found 288.4.

Step B: (4-Fluoro-phenyl)-(3-methyl-benzoylamino)-acetic acid **2** (34 mg, 0.12 mmol) was dissolved in DCM (2 mL), HOBt (20 mg, 0.14 mmol) and DIC (23 μL, 0.14 mmol) were added and the solution was stirred for 10 min at room temperature. N-(4-Methoxyphenyl)-ethane-1,2-diamine (24 mg, 0.14 mmol) was added and the solution was stirred for 3 h at

room temperature. The solvent was removed in vacuo, and the remainder was purified by reverse HPLC to afford N-{(4-Fluoro-phenyl)-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-methyl}-3-methyl-benzamide **3** (19 mg, 0.04 mmol, 36%) as a white solid: ¹H-NMR (400MHz, CD₃OD) δ = 7.62-6.83 (m, 12H), 5.51 (s, 1H), 3.79 (s, 3H), 3.77-3.46 (m, 4H), 2.32 (s, 3H). MS calcd. for C₂₅H₂₇FN₃O₃ (M+H⁺) 436.20, found 436.5.

[0328] Example 228. (S)-N-{2-(4-Hydroxy-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-O-*t*-butyl-tyrosine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (10 mg, 45%): ¹H-NMR (400MHz, CD₃OD) δ = 7.62-7.57 (m, 2H), 7.40-7.33 (m, 2H), 7.14-7.12 (m, 2H), 6.85-6.73 (m, 6H), 4.68 (t, *J*= 7.6, 1H), 3.75 (s, 3H), 3.41-3.38 (m, 2H), 3.25-3.01 (m, 4H), 2.41 (s, 3H). MS calcd. for C₂₆H₃₀N₃O₄ (M+H⁺) 448.22, found 448.2.

[0329] Example 229. (S)-N-{2-(2,4-Dichloro-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-2,4-dichloro phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (14 mg, 56%): MS calcd. for C₂₆H₂₈Cl₂N₃O₃ (M+H⁺) 500.15, found 500.1.

[0330] Example 230. (S)-N-{2-(3,5-Difluoro-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-3,5-difluoro phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (15 mg, 64%): MS calcd. for C₂₆H₂₈F₂N₃O₃ (M+H⁺) 468.21, found 468.2.

[0331] Example 231. (S)-N-{2-(3,4-Dichloro-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-3,4-dichloro phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (12 mg, 48%): MS calcd. for C₂₆H₂₈Cl₂N₃O₃ (M+H⁺) 500.15, found 500.1.

[0332] **Example 232.** (S)-4-Benzyloxy-N-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide; * C₃₂H₃₆F₃N₃O₄; ¹H NMR (CD₃Cl) δ(ppm) 7.61(m, 2H), 7.32(m, 5H), 6.96(m, 5H), 6.65(m, 2H), 6.42(m, 1H), 5.03(s, 2H), 4.46(dd, J=6.8Hz, J=15.2Hz, 1H), 3.44(m, 2H), 3.23(m, 2H), 1.59(m, 7H), 1.28(m, 1H), 1.11(m, 3H), 0.88(m, 2H); LCMS: 584.4(M+H)⁺.

[0333] **Example 233.** (S)-N-{2-(4-Acetylamino-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-4-acetylamino phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (13 mg, 53%): MS calcd. for C₂₈H₃₃N₄O₄ (M+H⁺) 489.25, found 489.2.

[0334] **Example 234.** (S)-Biphenyl-4-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₃₁H₃₄F₃N₃O₃; ¹H NMR (CD₃Cl) δ(ppm) 7.67(d, J=8.4, 2H), 7.51(d, J=8.4, 2H), 7.46(m, 3H), 7.27(m, 3H), 7.01(m, 2H), 6.89(d, J=8.8, 2H), 6.70(m, 1H), 4.41(m, 1H), 3.47(m, 2H), 3.28(m, 2H), 1.59(m, 7H), 1.28(m, 1H), 1.06(m, 3H), 0.84(m, 2H); LCMS: 554.4(M+H)⁺.

[0335] **Example 235.** (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-p-tolyl-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-4-methyl phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (12 mg, 54%): MS calcd. for C₂₇H₃₂N₃O₃ (M+H⁺) 446.24, found 446.3.

[0336] **Example 236.** (S)-N-{2-(3-Fluoro-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-3-fluoro phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (13 mg, 58%): MS calcd. for C₂₆H₂₉FN₃O₃ (M+H⁺) 450.22, found 450.2.

[0337] **Example 237.** (S)-N-{2-(3,4-Difluoro-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-3,4-difluoro phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was

prepared as a white solid (13 mg, 56%): MS calcd. for $C_{26}H_{28}F_2N_3O_3$ ($M+H^+$) 468.21, found 468.2.

[0338] **Example 238.** (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-m-tolyl-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting

5 Fmoc-(L)-phenyl alanine for Fmoc-(L)-3-methyl phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (13 mg, 58%): MS calcd. for $C_{27}H_{32}N_3O_3$ ($M+H^+$) 446.24, found 446.3.

[0339] **Example 239.** (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(2-trifluoromethyl-phenyl)-ethyl]-3-methyl-benzamide. * Following the procedure of

10 Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-2-trifluoromethyl phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (14 mg, 56%): MS calcd. for $C_{27}H_{29}F_3N_3O_3$ ($M+H^+$) 500.22, found 500.2.

[0340] **Example 240.** (S)-N-{2-(4-Cyano-phenyl)-1-[2-(4-methoxy-phenylamino)-

15 ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-4-cyano phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (14 mg, 61%): MS calcd. for $C_{27}H_{29}N_4O_3$ ($M+H^+$) 457.22 found 457.2.

[0341] **Example 241.** (S)-N-{2-(4-Bromo-phenyl)-1-[2-(4-methoxy-phenylamino)-

20 ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-4-bromo phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (12 mg, 47%): MS calcd. for $C_{26}H_{29}BrN_3O_3$ ($M+H^+$) 510.14, found 510.1.

[0342] **Example 242.** (S)-N-{2-(4-Iodo-phenyl)-1-[2-(4-methoxy-phenylamino)-

25 ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-4-iodo phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (13 mg, 47%): 1H -NMR (400MHz, CD_3OD) δ = 7.59-7.6.60 (m, 12H), 4.71 (dd, J = 7.0, J = 8.2, 1H), 3.69 (s, 3H), 3.38-2.97 (m, 6H), 2.37 (s, 3H). MS calcd. for $C_{26}H_{29}IN_3O_3$ ($M+H^+$) 558.12, found 558.1.

30

[0343] Example 243. (S)-N-{2-(4-Chloro-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-4-chloro phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (14 mg, 60%): MS calcd. for $C_{26}H_{29}ClN_3O_3$ ($M+H^+$) 466.19, found 466.2.

[0344] Example 244. (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-nitro-phenyl)-ethyl]-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-4-nitro phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (15 mg, 63%): MS calcd. for $C_{26}H_{29}N_4O_5$ ($M+H^+$) 477.21, found 477.2.

[0345] Example 245. (S)-N-{2-(4-Fluoro-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-4-fluoro phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (13 mg, 58%): MS calcd. for $C_{26}H_{29}FN_3O_3$ ($M+H^+$) 500.15, found 500.1.

[0346] Example 246. (S)-5-(3-Trifluoromethyl-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * $C_{30}H_{31}F_6N_3O_4$; 1H NMR (CD_3Cl) δ (ppm) 7.80(m, 2H), 7.46(m, 2H), 7.07(d, $J=4.0$ Hz, 1H), 6.87(m, 2H), 6.69(m, $J=3.6$ Hz, 1H), 6.60(m, 2H), 6.47(m, 2H), 4.48(m, 1H), 3.40(m, 2H), 3.16(m, 2H), 1.60(m, 7H), 1.26(m, 1H), 1.12(m, 3H), 0.86(m, 2H); LCMS: 612.4($M+H$) $^+$.

[0347] Example 247. (S)-N-{2-(4-Benzyloxy-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-4-benzyloxy phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (13 mg, 48%): MS calcd. for $C_{33}H_{35}N_3O_4$ ($M+H^+$) 538.27, found 538.3.

[0348] Example 248. (S)-N-{2-[4-(2,6-Dichloro-benzyloxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-4-(2,6-dichloro-benzyloxy) phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (11 mg, 36%): MS calcd. for $C_{33}H_{34}Cl_2N_3O_4$ ($M+H^+$) 606.19, found 606.2.

[0349] **Example 249.** (S)-N-{2-(4-Methoxy-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-4-methoxy phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (14 mg, 61%): MS calcd. for $C_{27}H_{32}N_3O_4$ ($M+H^+$) 462.24, found 462.2.

[0350] **Example 250.** 2-Amino-4-methyl-thiazole-5-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * $C_{23}H_{30}F_3N_5O_3S$; LCMS: 514.4($M+H$)⁺.

[0351] **Example 251.** (S)-5-(2-Chloro-5-trifluoromethyl-phenyl)-furan-2-carboxylic acid {2-cyclohexyl-1-[2-(4-trifluoromethoxy-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * $C_{30}H_{30}ClF_6N_3O_4$; ¹H NMR (CD_3Cl) δ (ppm) 7.92(m, 1H), 7.49(d, $J=8.52$ Hz, 1H), 7.42(m, 1H), 7.28(m, 1H), 7.08(d, $J=3.6$ Hz, 1H), 7.05(d, $J=4$ Hz, 1H), 7.00(d, $J=8.8$ Hz, 2H), 6.85(d, $J=8.8$ Hz, 2H), 6.78(m, 1H), 4.37(m, 1H), 3.48(m, 2H), 3.28(m, 2H), 1.64(m, 7H), 1.30(m, 1H), 1.06(m, 3H), 0.85(m, 2H); LCMS: 646.3($M+H$)⁺.

[0352] **Example 252.** (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(3-trifluoromethyl-phenyl)-ethyl]-3-methyl-benzamide. Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-3-trifluoromethyl phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (15 mg, 60%): MS calcd. for $C_{27}H_{29}F_3N_3O_3$ ($M+H^+$) 500.22, found 500.2.

[0353] **Example 253.** (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-trifluoromethyl-phenyl)-ethyl]-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-4-trifluoromethyl phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (15 mg, 60%): MS calcd. for $C_{27}H_{29}F_3N_3O_3$ ($M+H^+$) 500.22, found 500.2.

[0354] **Example 254.** (S)-N-{2-Benzyloxy-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-O-benzyl serine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (10 mg, 43%): MS calcd. for $C_{27}H_{32}N_3O_4$ ($M+H^+$) 462.24, found 462.2.

[0355] **Example 255.** (S)-N-{2-(4-tert-Butyl-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-4-tert-butyl phenyl alanine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (13 mg, 53%): MS calcd. for $C_{30}H_{38}N_3O_3$ ($M+H^+$) 488.29, found 488.3.

[0356] **Example 256.** 4-Cyclohexyl-N-[2-(4-methoxy-phenylamino)-ethyl]-2-(S)-(2-(S)-phenyl-propionylamino)-butyramide; * $C_{28}H_{39}N_3O_3$; LCMS: 466.6 ($M+H$)⁺.

[0357] **Example 257.** (S)-N-{2-(1H-Indol-3-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-N-Boc-tryptophane in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (8 mg, 34%): ¹H-NMR (400MHz, CD₃OD) δ = 7.69-6.71 (m, 13H), 4.80 (t, J =7.3, 1H), 3.72 (s, 3H), 3.43-3.30 (m, 4H), 3.15-3.02 (m, 2H), 2.40 (s, 3H). MS calcd. for $C_{28}H_{31}N_4O_3$ ($M+H^+$) 471.24, found 471.2.

[0358] **Example 258.** (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-naphthalen-1-yl-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(S)-2-amino-3-(1-naphthyl)-propionic acid in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (12 mg, 50%): MS calcd. for $C_{30}H_{32}N_3O_3$ ($M+H^+$) 482.24, found 482.2.

[0359] **Example 259.** (S)-N-{2-Benzoyloxy-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-propyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-O-benzyl threonine in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (8 mg, 34%): MS calcd. for $C_{28}H_{34}N_3O_4$ ($M+H^+$) 476.25, found 476.2.

[0360] **Example 260.** 3-Cyclohexyl-2-(S)-[2-(3-fluoro-phenyl)-acetylamino]-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide; * $C_{25}H_{31}F_2N_3O_2$; ¹H NMR (CDCl₃) δ (ppm) 8.60(m, 1H), 7.52(m, 2H), 7.40(m, 1H), 7.28(m, 2H), 7.11(m, 3H), 6.60(m, 1H), 4.28(m, 1H), 3.75(m, 5H), 3.55(m, 1H), 1.85(m, 6H), 1.70(m, 1H), 1.34(m, 4H), 1.05(m, 2H); LCMS: 444.5($M+H$)⁺.

[0361] **Example 261.** (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-naphthalen-2-yl-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(S)-2-amino-3-(2-naphtyl)-propionic acid in Step B and 3-methoxy benzoic acid for m-toluic acid in Step D, the title compound
5 was prepared as a white solid (13 mg, 54%): MS calcd. for $C_{30}H_{32}N_3O_3$ ($M+H^+$) 482.24, found 482.2.

[0362] **Example 262.** (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-pyridin-3-yl-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-3-pyridyl alanine in Step B and 3-
10 methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (3 mg, 14%): MS calcd. for $C_{25}H_{29}N_4O_3$ ($M+H^+$) 433.22, found 433.2.

[0363] **Example 263.** (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-pyridin-4-yl-ethyl}-3-methyl-benzamide. * Following the procedure of Preparation 1, except substituting Fmoc-(L)-phenyl alanine for Fmoc-(L)-4-pyridyl alanine in Step B and 3-
15 methoxy benzoic acid for m-toluic acid in Step D, the title compound was prepared as a white solid (5 mg, 23%): MS calcd. for $C_{25}H_{29}N_4O_3$ ($M+H^+$) 433.22, found 433.2.

[0364] **Example 264.** Furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * $C_{22}H_{22}FN_3O_3$; 1H NMR ($CDCl_3$) δ (ppm) 8.42(m, 1H), 7.41(m, 3H), 7.08(m, 3H), 6.89(m, 1H), 6.40(m, 1H), 4.30(m, 1H), 3.52(m,
20 4H), 1.66(m, 7H), 1.34(m, 1H), 1.12(m, 3H), 0.90(m, 2H); LCMS: 402.5 ($M+H$) $^+$.

[0365] **Example 265.** 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-(2-tetrazol-1-yl-acetyl-amino)-propionamide; * $C_{20}H_{28}FN_7O_2$; 1H NMR ($CDCl_3$) δ (ppm) 8.85(s, 1H), 8.32(m, 1H), 7.96(m, 1H), 7.31(m, 2H), 7.07(m, 2H), 5.28(d, $J=16.8$ Hz, 1H), 5.15(d, $J=16.8$ Hz, 1H), 4.28(m, 1H), 3.45(m, 4H), 1.52(m, 7H), 1.30(m, 4H), 0.82(m, 2H); LCMS:
25 418.5 ($M+H$) $^+$.

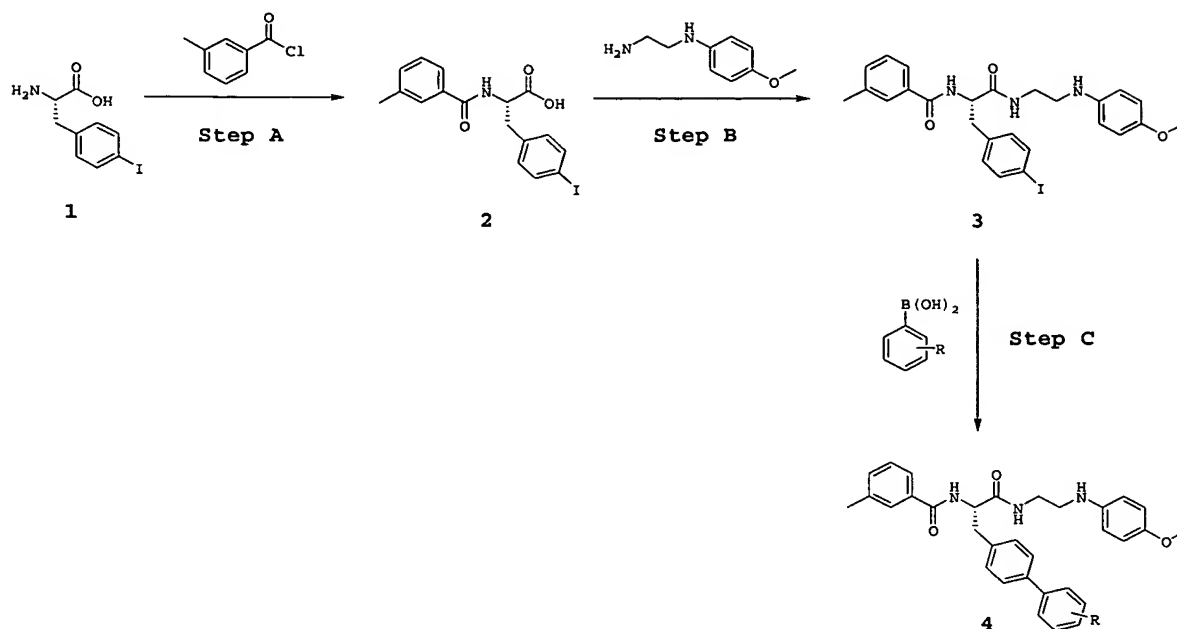
[0366] **Example 266.** N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-nitro-phenyl)-propyl]-3-methyl-benzamide. ** Following the procedure of Example 227, except substituting 4-fluoro-phenyl glycine for 2-amino-3-(4-nitro-phenyl)-butyric acid in Step A, the title compound was prepared as a white solid (7 mg, 29%): MS calcd. for $C_{27}H_{32}N_4O_5$
30 ($M+H^+$) 491.23, found 491.6.

[0367] **Example 267.** (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-m-tolyloxy-phenyl)-ethyl]-3-methyl-benzamide. ^{§§} Following the procedure of Example 327, except substituting phenyl-boronic acid for 3-methylphenyl-boronic acid in Step D, the title compound was prepared as a white solid (7 mg, 26%): ¹H-NMR (400MHz, CD₃OD) δ =
5 7.60-6.67 (m, 16H), 4.64 (dd, J = 7.1, J = 8.5, 1H), 3.81 (s, 3H), 3.60-3.07 (m, 6H), 2.37 (s, 3H), 2.26 (s, 3H). MS calcd. for C₃₃H₃₆N₃O₄ (M+H⁺) 538.27, found 538.4.

[0368] **Example 268.** threo-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-propyl}-3-methyl-benzamide. ^{**} Following the procedure of Example 227, except substituting 4-fluoro-phenyl glycine for threo-DL- β -methyl phenylalanine in Step A, the title
10 compound was prepared as a white solid (5 mg, 22%): ¹H-NMR (400MHz, CD₃OD) δ = 7.41-7.02 (m, 13H), 4.65 (d, J = 10.0, 1H), 3.81 (s, 3H), 3.77-2.96 (m, 5H), 2.30 (s, 3H), 1.33 (d, J = 7.0, 3H). MS calcd. for C₂₇H₃₂N₃O₃ (M+H⁺) 446.24, found 446.6.

[0369] **Example 269.** erythro-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-phenyl-propyl}-3-methyl-benzamide. ^{**} Following the procedure of Example 227, except
15 substituting 4-fluoro-phenyl glycine for erythro-DL- β -methyl phenylalanine in Step A, the title compound was prepared as a white solid (6 mg, 27%): ¹H-NMR (400MHz, CD₃OD) δ = 7.68-7.01 (m, 13H), 4.61 (d, J = 10.2, 1H), 3.80 (s, 3H), 3.77-2.96 (m, 5H), 2.40 (s, 3H), 1.41 (d, J = 6.9, 3H). MS calcd. for C₂₇H₃₂N₃O₃ (M+H⁺) 446.24, found 446.5.

[0370] **Example 270.** (S)-N-{2-Biphenyl-4-yl-1-[2-(4-methoxy-phenylamino)-ethyl]carbamoyl}-ethyl}-3-methyl-benzamide.



Step A: L-*p*-Iodo-phenylalanine (3.00 g, 10 mmol) was dissolved in H₂O (25 mL) containing
 5 equimolar amounts of NaOH (0.40 g, 10 mmol). The solution was cooled to 0 °C, then m-
 toluoyl chloride (1.32 mL, 10 mmol) was added dropwise under vigorous stirring. The
 mixture was allowed to warm to room temperature and stirred for approx. 2 h. After
 neutralization with 0.5 M HCl, the product was extracted from the reaction mixture three
 10 times with EtOAc. The combined organic layers were dried (MgSO₄), filtered and the
 solvent was removed in vacuo to yield (S)-(3-(4-Iodo-phenyl)-2-(3-methyl-benzoylamino)-
 propionic acid **2** (3.36 g, 82%): ¹H-NMR (400MHz, CD₃OD) δ = 7.62-7.05 (m, 8H), 4.82
 (dd, *J* = 5.0, *J* = 9.7, 1H), 3.32-3.26 (m, 1H), 3.08-3.03 (m, 1H), 2.36 (s, 3H). MS calcd. for
 C₁₇H₁₇INO₃ (M+H⁺) 410.22, found 410.2.

Step B: (S)-(3-(4-Iodo-phenyl)-2-(3-methyl-benzoylamino)-propionic acid **2** (3.36 g, 8.2
 15 mmol) was dissolved in DMF (40 mL), HOBt (1.20 g, 9 mmol) and DIC (1.38 ml, 8.9 mmol)
 were added and the solution was stirred for 10 min at room temperature. N-(4-
 Mthoxyphenyl)-ethane-1,2-diamine (1.48 g, 8.9 mmol) was added and the solution was
 stirred for 4 h at room temperature. The reaction mixture was then diluted with EtOAc and
 washed with H₂O three times. The organic solvent was partially removed, upon which the
 20 product precipitated. The precipitate was collected by filtration, recrystallized from EtOAc
 and dried under high vacuum to yield (S)-N-{2-(4-Iodo-phenyl)-1-[2-(4-methoxy-

phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide **3** (3.03 g, 68%): $^1\text{H-NMR}$ (400MHz, CD_3OD) δ = 7.59-7.6.60 (m, 12H), 4.71 (dd, J = 7.0, J = 8.2, 1H), 3.69 (s, 3H), 3.38-2.97 (m, 6H), 2.37 (s, 3H). MS calcd. for $\text{C}_{26}\text{H}_{29}\text{IN}_3\text{O}_3$ ($\text{M}+\text{H}^+$) 558.12, found 558.1.

Step C: (S)-N-{2-(4-Iodo-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-

5 methyl-benzamide **3** (20 mg, 0.04 mmol) was suspended in a 2:1 mixture of dioxane and water (3 mL) together with phenyl-boronic acid (13 mg, 0.12 mmol), Na_2CO_3 (15 mg, 0.16 mmol) and $\text{Pd}(\text{PPh}_3)_4$ (4 mg, 0.004 mmol). The mixture was heated to 150°C (microwave-assisted) for 5 min and filtered. The filtrate was subjected to reverse phase HPLC, the fractions containing the product were combined and lyophilized to yield the title compound
10 as a white solid (9 mg, 44%): $^1\text{H-NMR}$ (400MHz, CD_3OD) δ = 7.57-6.74 (m, 17H), 4.83 (m, 1H), 3.74 (s, 3H), 3.72-3.23 (m, 4H), 2.36 (s, 3H). MS calcd. for $\text{C}_{32}\text{H}_{34}\text{N}_3\text{O}_3$ ($\text{M}+\text{H}^+$) 508.26, found 508.2.

[0371] For the examples which were prepared according to the procedures in Example 270, partial or complete racemization at the stereogenic center of the α -amino acids may have
15 occurred.

[0372] **Example 271.** (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(3'-nitro-biphenyl-4-yl)-ethyl]-3-methyl-benzamide.^{##} Following the procedure of Example 270, except substituting phenyl-boronic acid for 3-nitrophenyl-boronic acid in Step C, the title compound was prepared as a white solid (8 mg, 36%): MS calcd. for $\text{C}_{32}\text{H}_{33}\text{N}_4\text{O}_5$ ($\text{M}+\text{H}^+$)
20 553.25, found 553.2.

[0373] **Example 272.** Furan-3-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide;* $\text{C}_{22}\text{H}_{28}\text{FN}_3\text{O}_3$; $^1\text{H NMR}$ (CDCl_3) δ (ppm) 8.48(m, 1H), 7.83(s, 1H), 7.40(m, 2H), 7.32(m, 1H), 7.07(m, 3H), 6.54(m, 1H), 4.31(m, 1H), 3.53(m, 3H), 3.40(m, 1H), 1.60(m, 7H), 1.30(m, 1H), 1.14(m, 3H), 0.87(m, 2H); LCMS:
25 402.5 ($\text{M}+\text{H}$)⁺.

[0374] **Example 273.** (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(2'-nitro-biphenyl-4-yl)-ethyl]-3-methyl-benzamide.^{##} Following the procedure of Example 270, except substituting phenyl-boronic acid for 2-nitrophenyl-boronic acid in Step C, the title compound was prepared as a white solid (8 mg, 36%): MS calcd. for $\text{C}_{32}\text{H}_{33}\text{N}_4\text{O}_5$ ($\text{M}+\text{H}^+$)
30 553.25, found 553.2.

[0375] **Example 274.** (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-pyridin-3-yl-phenyl)-ethyl]-3-methyl-benzamide.^{##} Following the procedure of Example

270, except substituting phenyl-boronic acid for 3-pyridyl-boronic acid in Step C, the title compound was prepared as a white solid (4 mg, 20%): MS calcd. for $C_{31}H_{33}N_4O_3$ ($M+H^+$) 509.26, found 509.2.

[0376] **Example 275.** (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-thiophen-3-yl-phenyl)-ethyl]-3-methyl-benzamide. ^{###} Following the procedure of Example 270, except substituting phenyl-boronic acid for 3-thienyl-boronic acid in Step C, the title compound was prepared as a white solid (9 mg, 44%): MS calcd. for $C_{30}H_{32}N_3O_3S$ ($M+H^+$) 514.22, found 514.2.

[0377] **Example 276.** (S)-N-{2-(4'-Cyano-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. ^{###} Following the procedure of Example 270, except substituting phenyl-boronic acid for 4-cyanophenyl-boronic acid in Step C, the title compound was prepared as a white solid (8 mg, 38%): MS calcd. for $C_{33}H_{33}N_4O_3$ ($M+H^+$) 533.26, found 533.2.

[0378] **Example 277.** (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-pyridin-4-yl-phenyl)-ethyl]-3-methyl-benzamide. ^{###} Following the procedure of Example 270, except substituting phenyl-boronic acid for 4-pyridyl-boronic acid in Step C, the title compound was prepared as a white solid (4 mg, 20%): MS calcd. for $C_{31}H_{33}N_4O_3$ ($M+H^+$) 509.26, found 509.2.

[0379] **Example 278.** (S)-N-{2-(4'-Chloro-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. ^{###} Following the procedure of Example 270, except substituting phenyl-boronic acid for 4-chlorophenyl-boronic acid in Step C, the title compound was prepared as a white solid (8 mg, 37%): MS calcd. for $C_{32}H_{33}ClN_3O_3$ ($M+H^+$) 542.22, found 542.2.

[0380] **Example 279.** (S)-N-{2-(2',3'-Dimethoxy-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. ^{###} Following the procedure of Example 270, except substituting phenyl-boronic acid for 2,3-dimethoxyphenyl-boronic acid in Step C, the title compound was prepared as a white solid (10 mg, 44%): MS calcd. for $C_{34}H_{38}N_3O_5$ ($M+H^+$) 568.28, found 568.3.

[0381] **Example 280.** (S)-N-{2-(3'-Amino-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. ^{###} Following the procedure of Example 270, except substituting phenyl-boronic acid for 3-aminophenyl-boronic acid in Step C, the title

compound was prepared as a white solid (8 mg, 38%): MS calcd. for $C_{32}H_{35}N_4O_3$ ($M+H^+$) 523.27, found 523.3.

[0382] **Example 281.** (S)-N-{2-(3',4'-Dimethoxy-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. ^{###} Following the procedure of
5 Example 270, except substituting phenyl-boronic acid for 3,4-dimethoxyphenyl-boronic acid in Step C, the title compound was prepared as a white solid (12 mg, 53%): MS calcd. for $C_{34}H_{38}N_3O_5$ ($M+H^+$) 568.28, found 568.3.

[0383] **Example 282.** (S)-N-{2-(4'-Hydroxymethyl-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. ^{###} Following the procedure of
10 Example 270, except substituting phenyl-boronic acid for 4-hydroxymethylphenyl-boronic acid in Step C, the title compound was prepared as a white solid (11 mg, 51%): MS calcd. for $C_{33}H_{36}N_3O_4$ ($M+H^+$) 538.27, found 538.2.

[0384] **Example 283.** (S)-N-{2-(5'-Fluoro-2'-methoxy-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. ^{###} Following the procedure of
15 Example 270, except substituting phenyl-boronic acid for 5-fluoro-2-methoxyphenyl-boronic acid in Step C, the title compound was prepared as a white solid (10 mg, 45%): MS calcd. for $C_{33}H_{35}FN_3O_4$ ($M+H^+$) 556.26, found 556.3.

[0385] **Example 284.** (S)-N-{2-(3'-Hydroxymethyl-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. ^{###} Following the procedure of
20 Example 270, except substituting phenyl-boronic acid for 3-hydroxymethylphenyl-boronic acid in Step C, the title compound was prepared as a white solid (11 mg, 51%): MS calcd. for $C_{33}H_{36}N_3O_4$ ($M+H^+$) 538.27, found 538.3.

[0386] **Example 285.** (S)-N-{2-(2',5'-Dimethoxy-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. ^{###} Following the procedure of
25 Example 270, except substituting phenyl-boronic acid for 2,5-dimethoxyphenyl-boronic acid in Step C, the title compound was prepared as a white solid (9 mg, 40%): MS calcd. for $C_{34}H_{38}N_3O_5$ ($M+H^+$) 568.28, found 568.3.

[0387] **Example 286.** (S)-N-{2-(2',5'-Dichloro-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. ^{###} Following the procedure of
30 Example 270, except substituting phenyl-boronic acid for 2,5-dichlorophenyl-boronic acid in

Step C, the title compound was prepared as a white solid (10 mg, 43%): MS calcd. for $C_{32}H_{32}Cl_2N_3O_3$ ($M+H^+$) 576.18, found 576.2.

[0388] Example 287. (S)-N-{2-(4'-Dimethylamino-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide.^{##} Following the procedure of Example 270, except substituting phenyl-boronic acid for 4-dimethylaminophenyl-boronic acid in Step C, the title compound was prepared as a white solid (5 mg, 23%): MS calcd. for $C_{34}H_{39}N_4O_3$ ($M+H^+$) 551.30, found 551.3.

[0389] Example 288. (S)-N-{2-(2'-Acetyl-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide.^{##} Following the procedure of Example 270, except substituting phenyl-boronic acid for 2-acetylphenyl-boronic acid in Step C, the title compound was prepared as a white solid (7 mg, 32%): MS calcd. for $C_{34}H_{36}N_3O_4$ ($M+H^+$) 550.27, found 550.3.

[0390] Example 289. (S)-N-{2-(4'-Hydroxy-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide.^{##} Following the procedure of Example 270, except substituting phenyl-boronic acid for 4-hydroxyphenyl-boronic acid in Step C, the title compound was prepared as a white solid (8 mg, 38%): MS calcd. for $C_{32}H_{34}N_3O_4$ ($M+H^+$) 524.25, found 524.3.

[0391] Example 290. (S)-N-{2-(3'-Acetyl-biphenyl-4-yl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide.^{##} Following the procedure of Example 270, except substituting phenyl-boronic acid for 3-acetylphenyl-boronic acid in Step C, the title compound was prepared as a white solid (12 mg, 55%): MS calcd. for $C_{34}H_{36}N_3O_4$ ($M+H^+$) 550.27, found 550.3.

[0392] Example 291. (S)-N-{2-[4-(2,4-Dimethoxy-pyrimidin-5-yl)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide.^{##} Following the procedure of Example 270, except substituting phenyl-boronic acid for 2,4-dimethoxy-5-pyrimidinyl-boronic acid in Step C, the title compound was prepared as a white solid (10 mg, 44%): MS calcd. for $C_{32}H_{36}N_5O_5$ ($M+H^+$) 570.27, found 570.3.

[0393] Example 292. (S)-N-{1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-[4-(6-methoxy-pyridin-3-yl)-phenyl]-ethyl}-3-methyl-benzamide.^{##} Following the procedure of Example 270, except substituting phenyl-boronic acid for 2-methoxy-5-pyridyl-boronic acid

in Step C, the title compound was prepared as a white solid (9 mg, 42%): MS calcd. for $C_{32}H_{35}N_4O_4$ ($M+H^+$) 539.27, found 539.3.

[0394] **Example 293.** 5-Methanesulfonyl-thiophene-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide; $C_{23}H_{30}FN_3O_4S_2$; 1H NMR ($CDCl_3$) δ (ppm) 8.58(m, 1H), 7.69(d, $J=5.2$, 1H), 7.54(d, $J=4.0$ Hz, 1H), 7.50(d, $J=4.0$ Hz, 1H), 7.44(m, 2H), 7.13(m, 2H), 4.34(m, 1H), 3.59(m, 3H), 3.38(m, 1H), 3.08(s, 3H), 1.66(m, 7H), 1.35(m, 1H), 1.13(m, 3H), 0.88(m, 2H); LCMS: 496.4 ($M+H$) $^+$.

[0395] **Example 294.** N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-2-(S)-phenyl-propionamide; $C_{26}H_{34}FN_3O_2$; 1H NMR ($CDCl_3$) δ (ppm) 7.62(m, 1H), 7.20(m, 5H), 6.95(m, 4H), 6.00(d, $J=5.6$ Hz, 1H), 4.07(m, 1H), 3.46(m, 3H), 3.23(m, 2H), 1.56(m, 6H), 1.42(m, 4H), 1.04(m, 4H), 0.81(m, 2H); LCMS: 440.5 ($M+H$) $^+$.

[0396] **Example 295.** Pyridazine-4-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide; $C_{22}H_{28}FN_5O_2$; 1H NMR ($CDCl_3$) δ (ppm) 9.62(m, 1H), 9.28(m, 1H), 8.59(d, $J=6.4$ Hz, 1H), 8.47(m, 1H), 8.12(m, 1H), 7.43(m, 2H), 7.08(m, 2H), 4.53(m, 1H), 3.53(m, 4H), 1.62(m, 7H), 1.36(m, 1H), 1.14(m, 3H), 0.90(m, 2H); LCMS: 414.5 ($M+H$) $^+$.

[0397] **Example 296.** N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-3-methanesulfonyl-benzamide; $C_{25}H_{32}FN_3O_4S$; 1H NMR ($CDCl_3$) δ (ppm) 8.57(m, 1H), 8.31(s, 1H), 7.95(m, 2H), 7.73(m, 1H), 7.49(m, 3H), 7.10(m, 2H), 4.42(m, 1H), 3.58(m, 3H), 3.45(m, 1H), 2.97(s, 3H), 1.62(m, 7H), 1.36(m, 1H), 1.14(m, 3H), 0.90(m, 2H); LCMS: 490.4 ($M+H$) $^+$.

[0398] **Example 297.** 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-(2-1H-tetrazol-5-yl-acetylamino)-propionamide; $C_{20}H_{28}FN_7O_2$; 1H NMR ($CDCl_3$) δ (ppm) 9.14(s, 1H), 8.62(m, 1H), 8.26(d, $J=5.6$ Hz, 1H), 7.59(m, 2H), 7.36(m, 2H), 5.58(d, $J=16.4$ Hz, 1H), 5.44(d, $J=16.4$ Hz, 1H), 4.56(m, 1H), 3.73(m, 4H), 1.81(m, 7H), 1.41(m, 4H), 1.12(m, 2H); LCMS: 418.5($M+H$) $^+$.

[0399] **Example 298.** Cyclopropanecarboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide; $C_{21}H_{30}FN_3O_2$; 1H NMR ($CDCl_3$) δ (ppm) 8.60(m, 1H), 7.52(m, 2H), 7.20(m, 2H), 6.78(d, $J=4$ Hz, 1H), 4.18(m, 1H), 3.69(m, 3H), 3.42(m, 1H), 1.74(m, 8H), 1.54(m, 1H), 1.26(m, 3H), 1.00(m, 2H), 0.85(m, 3H), 1.44(m, 1H); LCMS: 376.5($M+H$) $^+$.

[0400] **Example 299.** N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-4-methanesulfonylamino-benzamide; * C₂₅H₃₃FN₄O₄S; ¹H NMR (CDCl₃) δ(ppm) 8.68(s, 1H), 8.05(m, 1H), 7.61(m, 2H), 7.43(d, J=8.8Hz, 2H), 7.21(m, 1H), 7.17(m, 2H), 6.83(d, J=8.8Hz, 2H), 4.20(m, 1H), 3.82(m, 1H), 3.67(m, 1H), 3.43(m, 2H), 2.88(s, 3H), 1.74(m, 7H), 1.43(m, 1H), 1.20(m, 3H), 0.94(m, 2H); LCMS: 505.4(M+H)⁺.

[0401] **Example 300.** (S)-N-{2-[4-(4-Chloro-phenoxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide. ^{ss} Following the procedure of Example 327, except substituting phenyl-boronic acid for 4-chlorophenyl-boronic acid in Step D, the title compound was prepared as a white solid (4 mg, 14%): ¹H-NMR (400MHz, CD₃OD) δ = 7.60-6.86 (m, 16H), 4.66 (dd, J = 7.0, J = 8.6, 1H), 3.81 (s, 3H), 3.60-3.07 (m, 6H), 2.38 (s, 3H). MS calcd. for C₃₂H₃₃ClN₃O₄ (M+H)⁺ 558.22, found 558.4.

[0402] **Example 301.** 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-[2-(4-methoxy-phenyl)-acetylamino]-propionamide; * C₂₆H₃₄FN₃O₃; ¹H NMR (CDCl₃) δ(ppm) 8.26(m, 1H), 7.24(m, 2H), 7.08(m, 2H), 7.01(m, 2H), 6.75(m, 2H), 6.09(d, J=4.0Hz, 1H), 3.99(m, 1H), 3.69(s, 3H), 3.48(m, 5H), 3.28(m, 1H), 1.56(m, 6H), 1.42(m, 1H), 1.05(m, 4H), 0.79(m, 2H); LCMS: 456.5(M+H)⁺.

[0403] **Example 302.** 2-(S)-[2-(3-Chloro-phenyl)-acetylamino]-3-cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide; * C₂₅H₃₁FN₃O₂; ¹H NMR (CDCl₃) δ(ppm) 8.42(m, 1H), 7.34(m, 5H), 7.16(m, 3H), 6.47(d, J=4.4Hz, 1H), 4.17(m, 1H), 3.61(m, 5H), 3.41(m, 1H), 1.71(m, 6H), 1.59(m, 1H), 1.13(m, 4H), 0.86(m, 2H); LCMS: 460.5(³⁵ClM+H)⁺, 462.4(³⁷ClM+H)⁺.

[0404] **Example 303.** 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-phenylacetylamino-propionamide; * C₂₅H₃₂FN₃O₂; ¹H NMR (CDCl₃) δ(ppm) 8.37(m, 1H), 7.29(m, 6H), 7.24(m, 1H), 7.08(m, 2H), 6.23(d, J=4.0Hz, 1H), 4.09(m, 1H), 3.57(m, 5H), 3.37(m, 1H), 1.65(m, 6H), 1.59(m, 1H), 1.13(m, 4H), 0.86(m, 2H); LCMS: 426.4(M+H)⁺.

[0405] **Example 304.** 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-(2-o-tolyl-acetylamino)-propionamide; * C₂₆H₃₄FN₃O₂; ¹H NMR (CDCl₃) δ(ppm) 8.42(m, 1H), 7.34(m, 2H), 7.22(m, 4H), 7.09(m, 2H), 6.10(d, J=3.6Hz, 1H), 4.08(m, 1H), 3.57(m, 5H), 3.44(m, 1H), 2.25(s, 3H), 1.58(m, 7H), 1.11(m, 4H), 0.85(m, 2H); LCMS: 440.5(M+H)⁺.

[0406] **Example 305.** 2-(S)-[2-(4-Chloro-phenyl)-acetylamino]-3-cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-propionamide; * C₂₅H₃₁ClFN₃O₂; ¹H NMR (CDCl₃) δ(ppm)

8.33(m, 1H), 7.13(m, 8H), 6.29(d, $J=4.0$ Hz, 1H), 4.02(m, 1H), 3.48(m, 5H), 3.28(m, 1H), 1.62(m, 6H), 1.45(m, 1H), 1.12(m, 4H), 0.82(m, 2H); LCMS: 460.4($^{35}\text{ClM}+\text{H}$) $^+$, 462.5($^{37}\text{ClM}+\text{H}$) $^+$.

[0407] **Example 306.** 3-Cyclohexyl-2-(S)-[2-(2-fluoro-phenyl)-acetyl-amino]-N-[2-(4-fluoro-phenyl-amino)-ethyl]-propionamide; $^*\text{C}_{25}\text{H}_{31}\text{F}_2\text{N}_3\text{O}_2$; ^1H NMR (CDCl_3) δ (ppm) 8.20(m, 1H), 7.19(m, 3H), 7.16(m, 2H), 6.98(m, 4H), 6.29(d, $J=4.0$ Hz, 1H), 4.06(m, 1H), 3.48(m, 5H), 3.28(m, 1H), 1.61(m, 6H), 1.47(m, 1H), 1.08(m, 4H), 0.81(m, 2H); LCMS: 444.5(M+H) $^+$.

[0408] **Example 307.** 3-Cyclohexyl-2-(S)-diphenylacetyl-amino-N-[2-(4-fluoro-phenyl-amino)-ethyl]-propionamide; $^*\text{C}_{31}\text{H}_{36}\text{FN}_3\text{O}_2$; ^1H NMR (CDCl_3) δ (ppm) 8.46(m, 1H), 7.18(m, 10H), 6.91(m, 2H), 6.80(m, 2H), 6.20(d, $J=3.6$ Hz, 1H), 4.05(m, 1H), 3.51(m, 3H), 3.18(m, 1H), 1.50(m, 7H), 1.43(m, 1H), 1.01(m, 4H), 0.75(m, 2H); LCMS: 502.5(M+H) $^+$.

[0409] **Example 308.** N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenyl-amino)-ethyl-carbamoyl]-ethyl}-2-(2-fluoro-biphenyl-4-yl)-propionamide; $^*\text{C}_{32}\text{H}_{37}\text{F}_2\text{N}_3\text{O}_2$; ^1H NMR (CDCl_3) δ (ppm) 8.33(m, 1H), 7.35(m, 8H), 7.01(m, 4H), 6.30(m, 1H), 4.02(m, 1H), 3.60(m, 3H), 3.37(m, 2H), 1.52(m, 9H), 1.42(m, 1H), 1.04(m, 4H), 0.78(m, 2H); LCMS: 534.5(M+H) $^+$.

[0410] **Example 309.** N-{2-cyclohexyl-1-(S)-[2-(4-fluoro-phenyl-amino)-ethyl-carbamoyl]-ethyl}-2-p-tolyl-propionamide; $^*\text{C}_{27}\text{H}_{36}\text{FN}_3\text{O}_2$; ^1H NMR (CDCl_3) δ (ppm) 8.18(m, 1H), 7.37(m, 1H), 7.23(m, 1H), 7.02(m, 6H), 6.02(m, 1H), 3.95(m, 1H): 3.53(m, 3H), 3.31(m, 2H), 2.22(d, $J=15.2$ Hz, 3H), 1.55(m, 5H), 1.37(m, 5H), 1.01(m, 4H), 0.74(m, 2H); LCMS: 454.5(M+H) $^+$.

[0411] **Example 310.** N-{2-cyclohexyl-1-(S)-[2-(4-fluoro-phenyl-amino)-ethyl-carbamoyl]-ethyl}-2-(4-fluoro-phenyl)-propionamide; $^*\text{C}_{26}\text{H}_{33}\text{F}_2\text{N}_3\text{O}_2$; ^1H NMR (CDCl_3) δ (ppm) 8.25(m, 1H), 7.38(m, 1H), 7.21(m, 1H), 6.97(m, 6H), 6.18(m, 1H), 3.98(m, 1H), 3.55(m, 4H), 3.34(m, 1H), 1.58(m, 9H), 1.27(m, 1H), 1.06(m, 4H), 0.77(m, 2H); LCMS: 458.5 (M+H) $^+$.

[0412] **Example 311.** N-{2-cyclohexyl-1-(S)-[2-(4-fluoro-phenyl-amino)-ethyl-carbamoyl]-ethyl}-2-(4-hydroxy-phenyl)-propionamide; $^*\text{C}_{26}\text{H}_{34}\text{FN}_3\text{O}_3$; ^1H NMR (CDCl_3) δ (ppm) 8.18(m, 1H), 7.39(m, 1H), 7.26(m, 1H), 7.03(m, 4H), 6.68(m, 2H), 6.13(m, 1H), 3.96(m, 2H), 3.52(m, 4H), 3.31(m, 2H), 1.56(m, 8H), 1.18(m, 1H), 1.04(m, 4H), 0.74(m, 2H); LCMS: 465.5(M+H) $^+$.

[0413] **Example 312.** 2-(4-Chloro-phenyl)-N-{2-cyclohexyl-1-(S)-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-propionamide; * C₂₆H₃₃FN₃O₂; ¹H NMR (CDCl₃) δ(ppm) 8.24(m, 1H), 7.31(m, 1H), 7.16(m, 2H), 7.04(m, 5H), 6.13(m, 1H), 3.92(m, 1H), 3.50(m, 4H), 3.28(m, 1H), 1.54(m, 7H), 1.28(m, 3H), 0.99(m, 4H), 0.71(m, 2H); LCMS: 474.5(³⁵ClM+H)⁺, 476.5(³⁷ClM+H)⁺.

[0414] **Example 313.** N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-4-methanesulfonyl-benzamide; * C₂₅H₃₂FN₃O₄S; ¹H NMR (CDCl₃) δ(ppm) 8.55(m, 1H), 7.90(m, 4H), 7.52(m, 2H), 7.45(m, 1H), 7.19(m, 2H), 4.44(m, 1H), 3.68(m, 3H), 3.45(m, 1H), 3.04(s, 3H), 1.75(m, 7H), 1.46(m, 1H), 1.21(m, 3H), 0.99(m, 2H); LCMS: 490.4(M+H)⁺.

[0415] **Example 314.** Thiazole-4-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₁H₂₇FN₄O₂S; ¹H NMR (CDCl₃) δ(ppm) 8.97(m, 1H), 8.61(m, 1H), 8.15(m, 1H), 8.11(m, 1H), 7.70(m, 2H), 7.35(m, 2H), 4.56(m, 1H), 3.80(m, 4H), 1.96(m, 7H), 1.63(m, 1H), 1.39(m, 3H), 1.17(m, 2H); LCMS: 419.4(M+H)⁺.

[0416] **Example 315.** N-{2-cyclohexyl-1-(S)-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-2-(R)-phenyl-propionamide; * C₂₆H₃₄FN₃O₂; ¹H NMR (CDCl₃) δ(ppm) 8.46(m, 1H), 7.50(m, 2H), 7.34(m, 5H), 7.17(m, 2H), 6.16(m, 1H), 4.07(m, 1H), 3.65(m, 4H), 3.42(m, 1H), 1.47(m, 10H), 1.02(m, 4H), 0.76(m, 2H); LCMS: 440.5 (M+H)⁺.

[0417] **Example 316.** 4-Cyano-N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide; * C₂₅H₂₉FN₄O₂; ¹H NMR (CDCl₃) δ(ppm) 8.32(m, 1H), 7.64(d, J=8.4Hz, 2H), 7.47(d, J=8.4Hz, 2H), 7.38(d, J=6.0Hz, 1H), 7.25(m, 2H), 6.91(m, 2H), 4.26(m, 1H), 3.38(m, 3H), 3.26(m, 1H), 1.47(m, 7H), 1.18(m, 1H), 0.96(m, 3H), 0.76(m, 2H); LCMS: 437.5 (M+H)⁺.

[0418] **Example 317.** 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-(2-(R)-hydroxy-2-phenyl-acetyl-amino)-propionamide; * C₂₅H₃₂FN₃O₃; ¹H NMR (CDCl₃) δ(ppm) 8.36(m, 1H), 7.44(m, 2H), 7.28(m, 8H), 4.96(s, 1H), 4.12(m, 1H), 3.79(m, 2H), 3.45(m, 1H), 3.15(m, 1H), 1.75(m, 1H), 1.53(m, 6H), 0.89(m, 6H); LCMS: 442.5 (M+H)⁺.

[0419] **Example 318.** N-{2-cyclohexyl-1-(S)-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-2-(R)-phenyl-butyramide; * C₂₇H₃₆FN₃O₂; ¹H NMR (CDCl₃) δ(ppm) 8.02(m, 1H),

7.16(m, 7H), 6.98(m, 2H), 6.49(d, $J=5.6\text{Hz}$, 1H), 4.08(m, 1H), 3.45(m, 2H), 3.25(m, 3H), 1.94(m, 1H), 1.73(m, 1H), 1.55(m, 7H), 1.07(m, 4H), 0.79(m, 5H); LCMS: 454.5(M+H)⁺.

[0420] Example 319. 1-Phenyl-cyclopropanecarboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₇H₃₄FN₃O₂; ¹H NMR (CDCl₃)

5 δ(ppm) 8.31(m, 1H), 7.37(m, 4H), 7.27(m, 3H), 7.05(m, 2H), 5.77(d, $J=2.8\text{Hz}$, 1H), 3.82(m, 1H), 3.53(m, 3H), 3.29(m, 1H), 1.12(m, 14H), 0.80(m, 1H), 0.64(m, 2H); LCMS: 452.5(M+H)⁺.

[0421] Example 320. N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-2-(R,S)-(4-fluoro-phenyl)-propionamide; * C₂₆H₃₃F₂N₃O₂; ¹H NMR (CDCl₃) δ(ppm)

10 8.06(m, 1H), 7.22(m, 1H), 7.07(m, 1H), 6.89(m, 5H), 6.69(m, 2H), 3.82(m, 1H), 3.42(m, 4H), 3.20(m, 1H), 1.23(m, 9H), 1.16(m, 3H), 0.90(m, 3H), 0.66(m, 1H); LCMS: 496.5 (M+H)⁺.

[0422] Example 321. 3-Cyano-N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide; * C₂₅H₂₉FN₄O₂; ¹H NMR (CDCl₃) δ(ppm) 8.67(m, 1H),

15 8.04(m, 2H), 7.92(m, 3H), 7.62(m, 2H), 7.36(m, 2H), 4.65(m, 1H), 3.85(m, 4H), 1.90(m, 7H), 1.63(m, 1H), 1.37(m, 3H), 1.16(m, 2H); LCMS: 437.5(M+H)⁺.

[0423] Example 322. 5-(4-Fluoro-phenyl)-furan-2-carboxylic acid (S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-amide; * C₂₈H₃₁F₂N₃O₃; LCMS: 496.5 (M+H)⁺.

20 **[0424] Example 323.** 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-[2-(3-trifluoromethyl-phenyl)-acetyl-amino]-propionamide; * C₂₆H₃₁F₄N₃O₂; ¹H NMR (CDCl₃) δ(ppm) 8.26(m, 1H), 7.45(m, 4H), 7.24(m, 2H), 7.07(m, 2H), 6.39(d, $J=4.8\text{Hz}$, 1H), 4.17(m, 1H), 3.71(s, 2H), 3.57(m, 3H), 3.35(m, 1H), 1.68(m, 7H), 1.19(m, 4H), 0.91(m, 2H); LCMS: 494.5(M+H)⁺.

25 **[0425] Example 324.** 3-Cyano-N-(S)-{2-cyclohexyl-1-[2-(4-fluoro-phenylamino)-ethylcarbamoyl]-ethyl}-benzamide; * C₂₅H₂₉FN₄O₂; ¹H NMR (CDCl₃) δ(ppm) 8.09(m, 1H), 7.99(s, 1H), 7.91(m, 1H), 7.69(m, 1H), 7.46(m, 1H), 7.29(m, 1H), 7.24(m, 2H), 7.03(m, 2H), 4.42(m, 1H), 3.45(m, 3H), 3.37(m, 1H), 1.66(m, 7H), 1.34(m, 1H), 1.16(m, 3H), 0.90(m, 2H); LCMS: 437.5 (M+H)⁺.

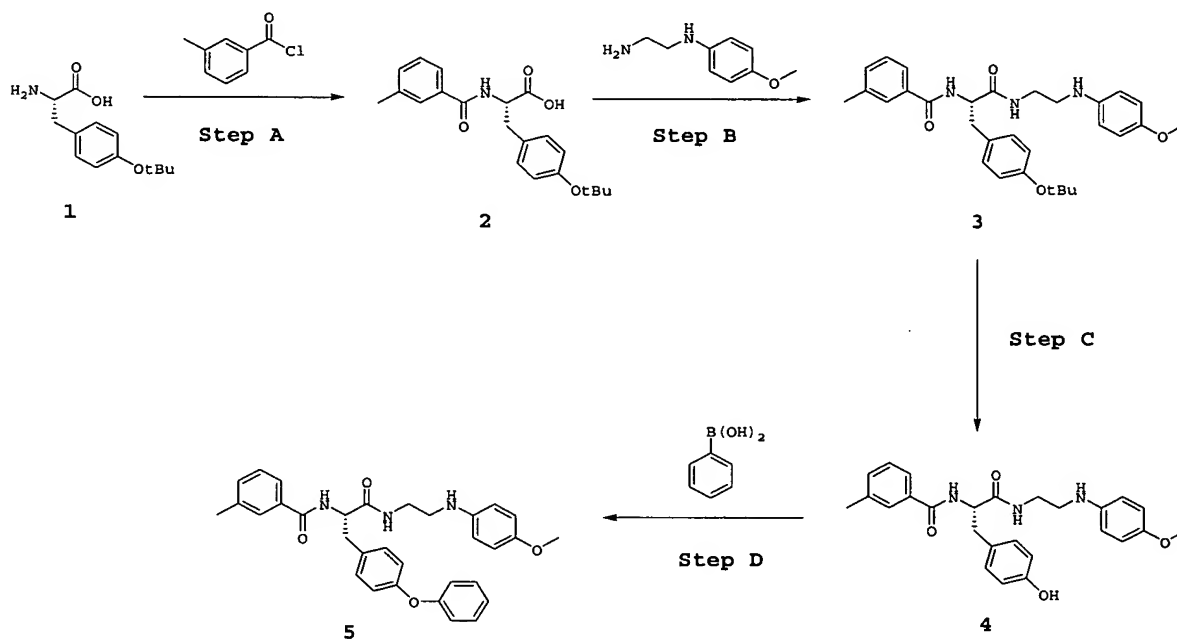
30 **[0426] Example 325.** 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-[2-(4-trifluoromethyl-phenyl)-acetyl-amino]-propionamide; * C₂₆H₃₁F₄N₃O₂; ¹H NMR (CDCl₃)

$\delta(\text{ppm})$ 8.57(M, 1H), 7.62(m, 2H), 7.43(m, 4H), 7.21(m, 2H), 6.63(m, 1H), 4.25(m, 1H), 3.85(m, 1H), 3.83(s, 2H), 3.72(m, 2H), 3.47(m, 1H), 1.73(m, 7H), 1.35(m, 4H), 1.05(m, 2H); LCMS: 494.5 (M+H)⁺.

[0427] **Example 326.** 3-Cyclohexyl-N-[2-(4-fluoro-phenylamino)-ethyl]-2-(S)-[2-(4-

methanesulfonyl-phenyl)-acetyl-amino]-propionamide; * C₂₆H₃₄FN₃O₄S; ¹H NMR (CDCl₃) $\delta(\text{ppm})$ 8.31(m, 1H), 7.72(d, *J*=8.4Hz, 2H), 7.36(d, *J*=8.4Hz, 2H), 7.24(m, 2H), 7.05(m, 2H), 6.60(m, 1H), 4.09(m, 1H), 3.65(s, 2H), 3.61(m, 1H), 3.50(m, 2H), 3.26(m, 1H), 2.93(s, 3H), 1.62(m, 7H), 1.12(m, 4H), 0.85(m, 2H); LCMS: 504.5(M+H)⁺.

[0428] **Example 327.** (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-phenoxy-phenyl)-ethyl]-3-methyl-benzamide.



Step A: O-t-Butyl-L-tyrosine **1** (2.00 g, 7.8 mmol) was dissolved in H₂O (10 mL) containing equimolar amounts of NaOH (0.31 g, 7.8 mmol). The solution was cooled to 0 °C, then m-toluoyl chloride (1.04 mL, 7.8 mmol) was added dropwise under vigorous stirring. The mixture was allowed to warm to room temperature and stirred for approx. 2 h. After acidification with 0.25 M phosphate buffer (pH 6.2), the product was extracted from the reaction mixture three times with EtOAc. The combined organic layers were dried (MgSO₄), filtered and the solvent was removed in vacuo to yield (S)-3-(4-tert-Butoxy-phenyl)-2-(3-methyl-benzoylamino)-propionic acid **2** (2.17 g, 6.1 mmol, 78%) as a white solid: ¹H-NMR (400MHz, CD₃OD) δ = 7.51-6.87 (m, 8H), 4.82 (dd, *J* = 5.0, *J* = 9.8, 1H), 3.33-3.28 (m, 1H),

3.09-3.03 (m, 1H), 2.35 (s, 3H), 1.27 (s, 9H). MS calcd. for $C_{21}H_{26}NO_4$ ($M+H^+$) 356.19, found 356.4.

Step B: (S)-3-(4-tert-Butoxy-phenyl)-2-(3-methyl-benzoylamino)-propionic acid **2** (0.80 g, 2.25 mmol) was dissolved in DCM (20 mL), HOBt (0.37 g, 2.7 mmol) and DIC (0.42 mL, 2.7 mmol) were added and the solution was stirred for 10 min at room temperature. N-(4-Methoxyphenyl)-ethane-1,2-diamine (0.45 g, 2.7 mmol) was added and the solution was stirred for 6 h at room temperature. The reaction mixture was then diluted with DCM and washed with H_2O three times. The organic layer was dried ($MgSO_4$), filtered and the solvent was removed in vacuo. The remainder was purified by chromatography (silica, DCM/MeOH gradient) and dried under high vacuum to afford (S)-N-{2-(4-tert-Butoxy-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide **3** (0.71 g, 1.4 mmol, 63%) as a white solid: 1H -NMR (400MHz, CD_3OD) δ = 7.55-6.79 (m, 12H), 4.80 (m, 1H), 3.76 (s, 3H), 3.52-3.47 (m, 2H), 3.24-3.15 (m, 4H), 2.37 (s, 3H), 1.32 (s, 9H). MS calcd. for $C_{30}H_{38}N_3O_4$ ($M+H^+$) 504.29, found 504.6.

Step C: (S)-N-{2-(4-tert-Butoxy-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide (0.50 g, 1.0 mmol) was dissolved in DCM containing 10% trifluoroacetic acid. The solution was stirred at room temperature for 1 h, then the solvents were removed in vacuo. The remaining crude material was purified by chromatography (silica, DCM/MeOH gradient) and dried under high vacuum to yield (S)-N-{2-(4-Hydroxy-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide **4** (0.40 g, 0.9 mmol, 90%): 1H -NMR (400MHz, CD_3OD) δ = 7.62-7.57 (m, 2H), 7.40-7.33 (m, 2H), 7.14-7.12 (m, 2H), 6.85-6.73 (m, 6H), 4.68 (t, J = 7.6, 1H), 3.75 (s, 3H), 3.41-3.38 (m, 2H), 3.25-3.01 (m, 4H), 2.41 (s, 3H). MS calcd. for $C_{26}H_{30}N_3O_4$ ($M+H^+$) 448.22, found 448.2.

Step D: (S)-N-{2-(4-Hydroxy-phenyl)-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide (0.02 g, 0.05 mmol) **4** was suspended in DCM (1 mL) together with phenyl-boronic acid (7 mg, 0.06 mmol), $Cu(OAc)_2$ (16 mg, 0.09 mmol) and NEt_3 (30 μL , 0.23 mmol). The mixture was stirred at room temperature for 14 h. The solvent was removed and the crude products were separated on reversed phase HPLC. The fraction containing the title compound **5** (S)-N-[1-[2-(4-Methoxy-phenylamino)-ethylcarbamoyl]-2-(4-phenoxy-phenyl)-ethyl]-3-methyl-benzamide ($R = H$) was lyophilized to yield a white solid (5 mg, 0.009 mmol, 19%): 1H -NMR (400MHz, CD_3OD) δ = 7.60-6.89 (m, 17H), 4.65 (dd, J = 7.1, J = 8.4, 1H), 3.81 (s, 3H), 3.61-3.07 (m, 6H), 2.37 (s, 3H). MS calcd. for $C_{32}H_{34}N_3O_4$ ($M+H^+$) 524.25, found 524.4.

[0429] For the examples which were prepared according to the procedures in Example 327, partial or complete racemization at the stereogenic center of the α -amino acids may have occurred.

[0430] **Example 328.** (S)-N-{2-[4-(4-Methoxy-phenoxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide.^{ss} Following the procedure of Example 327, except substituting phenyl-boronic acid for 4-methoxyphenyl-boronic acid in Step D, the title compound was prepared as a white solid (6 mg, 22%): ¹H-NMR (400MHz, CD₃OD) δ = 7.60-6.82 (m, 16H), 4.62 (dd, J = 7.2, J = 8.3, 1H), 3.81 (s, 3H), 3.77 (s, 3H), 3.61-3.05 (m, 6H), 2.38 (s, 3H). MS calcd. for C₃₃H₃₆N₃O₅ (M+H⁺) 554.27, found 554.4.

[0431] **Example 329.** (S)-N-{2-[4-(3-Chloro-phenoxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide.^{ss} Following the procedure of Example 327, except substituting phenyl-boronic acid for 3-chlorophenyl-boronic acid in Step D, the title compound was prepared as a white solid (4 mg, 14%): ¹H-NMR (400MHz, CD₃OD) δ = 7.61-6.82 (m, 16H), 4.66 (dd, J = 7.1, J = 8.5, 1H), 3.81 (s, 3H), 3.61-3.09 (m, 6H), 2.38 (s, 3H). MS calcd. for C₃₂H₃₃ClN₃O₄ (M+H⁺) 558.22, found 558.4.

[0432] **Example 330.** (S)-N-{2-[4-(3,5-Dimethyl-phenoxy)-phenyl]-1-[2-(4-methoxy-phenylamino)-ethylcarbamoyl]-ethyl}-3-methyl-benzamide.^{ss} Following the procedure of Example 327, except substituting phenyl-boronic acid for 3,5-dimethylphenyl-boronic acid in Step D, the title compound was prepared as a white solid (6 mg, 22%): ¹H-NMR (400MHz, CD₃OD) δ = 7.52-6.43 (m, 15H), 4.56 (dd, J = 7.1, J = 8.5, 1H), 3.73 (s, 3H), 3.55-2.99 (m, 6H), 2.29 (s, 3H), 2.13 (s, 6H). MS calcd. for C₃₄H₃₈N₃O₄ (M+H⁺) 552.29, found 552.4.

B. Assays for Cathepsin Inhibitory Activity

Cathepsin S

[0433] The optimal substrate for cathepsin S, acetyl-histidine-proline-valine-lysine-amino carbamoyl coumarin, was identified from screening a combinatorial library of fluorogenic peptide substrates (Harris, J. L., B. J. Backes, *et al.*, *Proc Natl Acad Sci U S A* **2000**, 97(14), 7754-9). Kinetic measurements are performed in a total reaction volume of 30 μ l in 384-well microtiter plates. Cathepsin S, at a final concentration of 0.3-3 nM (active site), is incubated with the compounds at twelve varying concentrations in a buffer containing 100mM NaAc (pH5.5), 1mM EDTA, 100mM NaCl, 0.01% Brij-35 for 20 minutes at room temperature. Control reactions in the absence of inhibitor are performed in replicates of 24. The reactions

are initiated by adding the substrate, acetyl-histidine-proline-valine-lysine-amino carbamoyl coumarin, to a final concentration of 50 μ M. The rate of substrate hydrolysis is measured by monitoring the increase in fluorescence at an excitation wavelength of 380nm and an emission wavelength of 450 nm that results from cleavage of the aniline bond in the substrate by the enzyme. The apparent inhibition constants for the compounds are determined from the enzyme progress curves (Kuzmic, P., K. C. Elrod, *et al.*, *Anal Biochem* **2000**, 286(1), 45-50) and are then used to calculate the inhibition constants for competitive inhibitors.

Cathepsin K

[0434] The optimal substrate for cathepsin K, acetyl-lysine-histidine-proline-lysine-amino carbamoyl coumarin, was identified from screening a combinatorial library of fluorogenic peptide substrates (Harris, J. L., B. J. Backes, *et al.*, *Proc Natl Acad Sci U S A* **2000**, 97(14), 7754-9). Kinetic measurements are performed in a total reaction volume of 30 μ l in 384-well microtiter plates. Cathepsin K, at a final concentration of 3.5 nM (active site), is incubated with the compounds at twelve varying concentrations in a buffer containing 100mM NaAc (pH5.5), 1mM EDTA, 100mM NaCl, 0.01% Brij-35 for 20 minutes at room temperature. Control reactions in the absence of inhibitor are performed in replicates of 24. The reactions are initiated by adding the substrate, acetyl-lysine-histidine-proline-lysine-amino carbamoyl coumarin, to a final concentration of 40 μ M. The rate of substrate hydrolysis is measured by monitoring the increase in fluorescence at an excitation wavelength of 380nm and an emission wavelength of 450 nm that results from cleavage of the aniline bond in the substrate by the enzyme. The apparent inhibition constants for the compounds are determined from the enzyme progress curves (Kuzmic, P., K. C. Elrod, *et al.*, *Anal Biochem* **2000**, 286(1), 45-50) and are then used to calculate the inhibition constants for competitive inhibitors.

Cathepsin L

[0435] The optimal substrate for cathepsin L, acetyl-histidine-lysine-phenylalanine-lysine-amino carbamoyl coumarin, was identified from screening a combinatorial library of fluorogenic peptide substrates (Harris, J. L., B. J. Backes, *et al.*, *Proc Natl Acad Sci U S A* **2000**, 97(14), 7754-9). Kinetic measurements are performed in a total reaction volume of 30 μ l in 384-well microtiter plates. Cathepsin L, at a final concentration of 0.1 nM (active site), is incubated with the compounds at twelve varying concentrations in a buffer containing 100mM NaAc (pH5.5), 1mM EDTA, 100mM NaCl, 0.01% Brij-35 for 20 minutes at room temperature. Control reactions in the absence of inhibitor are performed in replicates of 24. The reactions are initiated by adding the substrate, acetyl-histidine-lysine-phenylalanine-

lysine-amino carbamoyl coumarin, to a final concentration of 20 μM . The rate of substrate hydrolysis is measured by monitoring the increase in fluorescence at an excitation wavelength of 380nm and an emission wavelength of 450 nm that results from cleavage of the aniline bond in the substrate by the enzyme. The apparent inhibition constants for the compounds are determined from the enzyme progress curves (Kuzmic, P., K. C. Elrod, *et al.*, *Anal Biochem* **2000**, 286(1), 45-50) and are then used to calculate the inhibition constants for competitive inhibitors.

Cathepsin B

[0436] The optimal substrate for cathepsin B, acetyl-histidine-proline-valine-lysine-amino carbamoyl coumarin, was identified from screening a combinatorial library of fluorogenic peptide substrates (Harris, J. L., B. J. Backes, *et al.*, *Proc Natl Acad Sci U S A* **2000**, 97(14), 7754-9). Kinetic measurements are performed in a total reaction volume of 30 μl in 384-well microtiter plates. Cathepsin B, at a final concentration of 1.5 nM (active site), is incubated with the compounds at twelve varying concentrations in a buffer containing 100mM NaAc (pH5.5), 1mM EDTA, 100mM NaCl, 0.01% Brij-35 for 20 minutes at room temperature. Control reactions in the absence of inhibitor are performed in replicates of 24. The reactions are initiated by adding the substrate, acetyl-histidine-proline-valine-lysine-amino carbamoyl coumarin, to a final concentration of 10 μM . The rate of substrate hydrolysis is measured by monitoring the increase in fluorescence at an excitation wavelength of 380nm and an emission wavelength of 450 nm that results from cleavage of the aniline bond in the substrate by the enzyme. The apparent inhibition constants for the compounds are determined from the enzyme progress curves (Kuzmic, P., K. C. Elrod, *et al.*, *Anal Biochem* **2000**, 286(1), 45-50) and are then used to calculate the inhibition constants for competitive inhibitors.

[0437] Preferred cathepsin S inhibition constants for compounds of the present invention are less than 10 μM . More preferred inhibition constants for compounds of the present invention are less than 1.0 μM . Most preferred inhibition constants for compounds of the present invention are less than 0.1 μM .

[0438] Selectivity for cathepsin S in the presence of cathepsin isozymes was determined by the ratio of the cathepsin isozyme inhibition constant of a compound of the present invention to the cathepsin S inhibition constant of the same compound. Preferred compounds of the present invention selective for cathepsin S have ratios of greater than 10. More preferred compounds of the present invention selective for cathepsin S have ratios of greater than 100.

Most preferred compounds of the present invention selective for cathepsin S have ratios of greater than 1000.

Table II: Assay Data for Inhibitors of Cathepsin S

5

Compound	K _i Cat. S ^a	Selectivity for Cat. S over Cat. K ^b
1	++	++
2	+++	+
3	+++	+++
4	+++	+++
5	++	+
6	+++	++
7	++	++
8	+++	+++
9	++	+
10	+++	+++
11	+++	+
12	+++	+
13	+++	+++
14	++	++
15	++	+
16	++	+
17	+++	++
18	+	+
19	++	+
20	++	++
21	+++	+
22	++	+
23	+++	++
24	+++	+
25	+++	++
26	+++	++
27	+++	+++
28	++	+
29	+++	++
30	+++	+
31	+++	+
32	+++	+
33	+	+
34	+++	+++
35	+++	+
36	+++	++
37	+++	++
38	+++	++
39	++	++
40	++	+
41	++	++
42	++	+
43	+++	+
44	++	++

Compound	K _i Cat. S ^a	Selectivity for Cat. S over Cat. K ^b
45	+++	+++
46	++	++
47	+++	+++
48	++	++
49	+++	++
50	+++	+++
51	+++	+++
52	+	+
53	+++	+++
54	+++	+
55	++	++
56	++	++
57	++	+
58	+++	++
59	++	++
60	+++	+
61	++	+
62	+++	++
63	+	+
64	+	+
65	+++	++
66	++	++
67	+++	+
68	+++	+++
69	+++	+
70	+++	+++
71	+++	++
72	+	++
73	+++	++
74	+++	+++
75	+++	+++
76	+++	+++
77	+++	++
78	+++	+
79	+	+
80	++	+++
81	++	++
82	++	+
83	+++	++
84	+++	++
85	+++	++
86	+++	++
87	+++	++
88	++	+
89	++	+
90	++	+
91	+	+
92	+++	+
93	++	+++
94	+	+
95	+++	++
96	+	+

Compound	K_i Cat. S ^a	Selectivity for Cat. S over Cat. K ^b
97	++	+
98	++	++
99	+++	++
100	+++	+++
101	++	+
102	+++	++
103	++	++
104	+	+
105	+++	+
106	+++	++
107	+++	+++
108	+++	++
109	+++	+++
110	+++	++
111	+++	++
112	+++	++
113	+++	+
114	+++	++
115	+++	+++
116	+++	+++
117	+++	++
118	+++	++
119	+++	+
120	+++	++
121	+++	+
122	+++	+++
123	+++	++
124	+++	+
125	+++	++
126	+++	+++
127	++	++
128	+++	+++
129	+	+
130	+	+
131	+	+
132	++	++
133	++	++
134	++	++
135	+++	+++
136	++	++
137	+++	+++
138	+++	+++
139	+++	+++
140	+++	+++
141	+++	+
142	+++	+++
143	++	++
144	++	+++
145	+++	+++
146	+++	+++
147	+++	+++
148	+++	+++

Compound	K_i Cat. S ^a	Selectivity for Cat. S over Cat. K ^b
149	+++	+++
150	+++	+
151	+++	+
152	+++	++
153	+++	+
154	+++	++
155	+++	+++
156	+++	+++
157	+++	+++
158	+++	++
159	+++	+++
160	+	++
161	+++	++
162	+++	+
163	+++	+
164	+++	+++
165	+++	+++
166	+++	+++
167	+++	+++
168	+++	+++
169	++	++
170	+	+
171	+++	+++
172	+++	+++
173	+++	+++
174	+++	+
175	+++	+++
176	+++	+++
177	+++	++
178	+++	+++
179	+++	+++
180	+++	+
181	+++	+++
182	+++	+
183	+++	+++
184	+++	+++
185	+++	++
186	+++	++
187	+++	+
188	+++	+++
189	+++	++
190	+++	+++
191	+++	+
192	+++	+++
193	+++	++
194	++	+
195	++	+++
196	+++	++
197	+++	+++
198	++	+
199	++	++
200	+++	+

Compound	K_i Cat. S ^a	Selectivity for Cat. S over Cat. K ^b
201	+++	++
202	+++	+++
203	+++	+++
204	++	++
205	+++	++
206	++	++
207	++	++
208	+++	+++
209	+++	+++
210	+++	++
211	++	++
212	+++	++
213	+++	+++
214	+++	+++
215	+++	+++
216	++	+++
217	++	+++
218	++	+++
219	+++	++
220	++	++
221	+++	+
222	+++	+++
223	++	++
224	++	++
225	+	+
226	+	+
227	++	++
228	+++	+++
229	+++	+++
230	+++	+++
231	+++	+++
232	+++	+
233	+++	+++
234	+++	+
235	+++	+++
236	+++	+++
237	+++	+++
238	+++	+
239	+++	+++
240	+++	+++
241	+++	+++
242	+++	+++
243	+++	+++
244	++	+++
245	+++	+++
246	+++	++
247	+++	+++
248	+++	+++
249	+++	+++
250	+++	++
251	+++	+++
252	+++	+++

Compound	K_i Cat. S ^a	Selectivity for Cat. S over Cat. K ^b
253	+++	+++
254	++	++
255	++	++
256	+++	+++
257	+++	+++
258	+++	+++
259	++	++
260	+++	++
261	+++	+++
262	+	+
263	+	+
264	+++	++
265	++	++
266	++	+
267	+++	+++
268	++	++
269	+	+
270	+++	+++
271	++	++
272	+++	++
273	++	++
274	+++	+++
275	+++	+++
276	+++	+++
277	+++	++
278	++	++
279	++	++
280	++	++
281	++	++
282	++	++
283	++	++
284	++	++
285	++	++
286	+	+
287	++	++
288	++	++
289	++	++
290	+++	+++
291	++	++
292	+++	+++
293	+++	++
294	+++	++
295	++	++
296	+++	+++
297	++	++
298	+++	++
299	+++	+
300	+++	++
301	+++	++
302	+++	++
303	+++	++
304	+++	+++

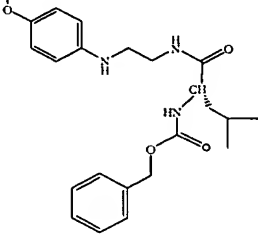
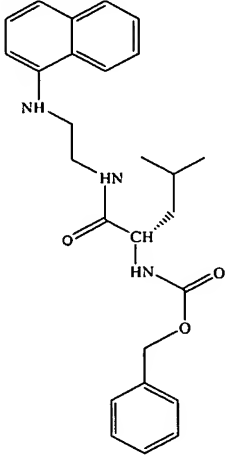
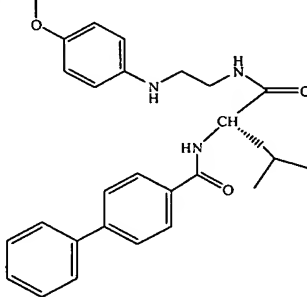
Compound	K _i Cat. S ^a	Selectivity for Cat. S over Cat. K ^b
305	+++	++
306	+++	+++
307	+++	+++
308	+++	++
309	+++	++
310	+++	++
311	+++	+
312	+++	++
313	++	+
314	+++	+
315	+++	++
316	+++	++
317	+++	++
318	+++	+
319	+++	++
320	+++	++
321	++	++
322	+++	++
323	+++	++
324	+++	++
325	+++	++
326	++	++
327	++	++
328	++	++
329	++	++
330	++	++

^a Cathepsin S inhibition constant for compounds of Formula I: +, <10 μ M; ++, <1.0 μ M; +++, <0.1 μ M.

^b Selectivity of compounds of Formula I for cathepsin S over cathepsin K: +, >10; ++, >100; +++, >1000.

C. Comparison Activity

- 5 [0439] In order to show the superiority of the compounds of the present invention versus compounds in the art, several compounds were tested in assays discussed herein. The compounds of the present invention showed superior unexpected properties, especially with respect to Cat S inhibition. Moreover, the compounds of the present invention were also more selective in inhibiting Cat S over Cat K.

Compound	Ki (CatS) ^a μM	Ki (CatK) ^b μM	Selectivity for Cat. S over Cat. K ^c
	++	+++	<10
	++	+++	<10
	+++	+++	<10

^a Cathepsin S inhibition constant for the compounds +, <10 μM; ++, <1.0 μM; +++, <0.1 μM.

^a Cathepsin K inhibition constant for the compounds +, <10 μM; ++, <1.0 μM; +++, <0.1 μM.

^c Selectivity of the compounds for cathepsin S over cathepsin K: <10

- 5 [0440] Although the foregoing invention has been de scribed in some detail by way of illustration and example for purposes of clarity of understanding, it will be obvious that certain changes and modifications may be practiced within the scope of the appended claims. In addition, each reference provided herein is incorporated by reference in its entirety to the same extent as if each reference was individually incorporated by reference.